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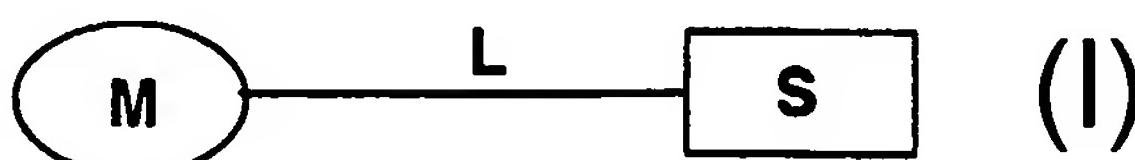
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- (71) Applicant (*for all designated States except US*): PLIVA D.D. [HR/HR]; Ulica grada Vukovara 49, 10000 Zagreb (HR). For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.
- (72) Inventors; and (75) Inventors/Applicants (*for US only*): MERCEP, Mladen [HR/HR]; Majstora Rádonje 10, 10000 Zagreb (HR). MESIC, Milan [HR/HR]; Slavenskog 8, 10000 Zagreb (HR). TOMASKOVIC, Linda [HR/HR]; Lascinska 119, 10000 Zagreb (HR). MARKOVIC, Stribor [HR/HR]; Sarajevska 2A, 47000 Karlovac (HR). MAKARUHA, Oresta [HR/HR]; Srebrnjak 5, 10000 Zagreb (HR). POLJAK, Visnja [HR/HR]; Poljica 29, 23000 Zadar (HR).

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(54) Title: NEW COMPOUNDS, COMPOSITIONS AND METHODS FOR TREATMENT OF INFLAMMATORY DISEASES AND CONDITIONS



steroid drug with anti-inflammatory activity and L represents a linker molecule linking M and S, (b) to their pharmacologically acceptable salts, prodrugs and solvates, (c) to processes and intermediates for their preparation, and (d) to their use in the treatment of inflammatory diseases and conditions in humans and animals. Such compounds inhibit many cytokines and immune mediators involved in immune responses which cause inflammation, allergy, or alloimmunity, including without limitation IL-1, 2, 4, 5, 6, 10, 12, GMCSF, ICAM, and TNF- α . Importantly, anti-inflammatory steroids exert a direct anti-inflammatory effect through binding to the glucocorticosteroid receptor.

(57) Abstract: The present invention relates (a) to new compounds represented by Formula I: wherein M represents a macrolide subunit (macrolide moiety) derived from macrolide possessing the property of accumulation in inflammatory cells, S represents a steroid subunit (steroid moiety) derived from

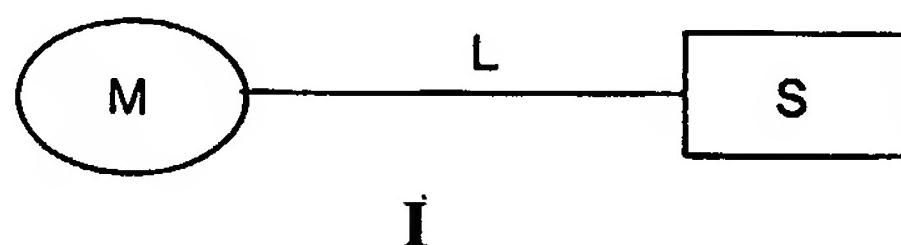
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NEW COMPOUNDS, COMPOSITIONS AND METHODS FOR TREATMENT OF INFLAMMATORY DISEASES AND CONDITIONS

This application claims priority to U.S. Provisional Application 60/394,670 filed July 8, 2002.

TECHNICAL PROBLEM

The present invention relates (a) to new compounds represented by the structure I:



wherein M represents a macrolide subunit possessing the property of accumulation in inflammatory cells, S represents an anti-inflammatory steroid subunit, and L represents a linker molecule linking M and S, (b) to their pharmacologically acceptable salts, prodrugs and solvates, (c) to processes and intermediates for their preparation, and (d) to their use in the treatment of inflammatory diseases and conditions in humans and animals. Such compounds inhibit many cytokines and immune mediators involved in immune responses which cause inflammation, allergy, or alloimmunity, including without limitation IL-1, 2, 4, 5, 6, 10, 12, GMCSF, ICAM, and TNF- α . Importantly, anti-inflammatory steroids exert a direct anti-inflammatory effect through binding to the glucocorticoid receptor.

PRIOR ART

Anti-inflammatory medicaments could be classified into those of steroid and of nonsteroidal type. Steroid anti-inflammatory compounds are still the most effective ones in the treatment of inflammatory diseases and conditions such as: asthma, chronic obstructive pulmonary disease, inflammatory nasal diseases such as allergic

rhinitis, nasal polyps, intestinal diseases such as Crohn's disease, colitis, ulcerative colitis, dermatological inflammations such as eczema, psoriasis, allergic dermatitis, neurodermatitis, pruritus, conjunctivitis, autoimmune diseases such as rheumatoid arthritis, and inhibition of transplantation immunity. Moreover, steroids are used as adjunct chemotherapeutic agents in treating various malignancies, including leukemias, lymphomas, myelomas, and other malignancies of the hematopoietic system. In addition to excellent potency and effectiveness, medicaments of this type also possess numerous unfavourable side-effects, e.g., on carbohydrate metabolism, calcium resorption, secretion of endogenous corticosteroids as well as on the physiological functions of the pituitary gland, adrenal cortex and thymus. Recently developed steroids are highly effective against inflammatory conditions and processes since they inhibit many inflammation mediators, whereas their systemic side-effects are diminished. Patent applications WO 94/13690, WO 94/14834, WO 92/13873 and WO 92/13872 describe the so-called "soft" steroids or hydrolysable corticosteroids designed for topical application on the inflammation site, whereas their systemic side-effects are diminished due to instability of "soft" steroids in the serum, wherein the active steroid very rapidly hydrolyses into the inactive form. An ideal steroid, however, without unfavourable effects in a long-term and continuous treatment as required for the control of diseases such as asthma or Crohn's disease has yet to be found. Thus there is an acute need for steroids with an improved therapeutic profile, and/or fewer or milder side effects.

Macrolides such as macrolide antibiotics accumulate preferentially within different cells of subjects administered such molecules, especially within phagocyte cells such as mononuclear peripheral blood cells, peritoneal and alveolar macrophages as well as in the liquid surrounding the bronchoalveolar epithelium (Glaude R. P. et al., *Antimicrob. Agents Chemother.*, 33 1989, 277-282; Olsen K. M. et al., *Antimicrob. Agents Chemother.*, 40 1996, 2582-2585). Moreover, relatively weak inflammatory effects of some macrolides have been described. For example, the anti-inflammatory effect of erythromycin derivatives (*J. Antimicrob. Chemother.*, 41 1998 37-46; WO 00/42055) and azithromycin derivatives has recently been described (EP 0283055). Anti-inflammatory effects of some macrolides are also known from *in vitro* and *in vivo* studies in experimental animal models such as zimosane-induced

peritonitis in mice (*J. Antimicrob. Chemother.*, 30 1992 339-348) and endotoxin-induced neutrophil accumulation in rat trachea (*J. Immunol.* 159 1997 3395-4005). The modulating effect of macrolides upon cytokines such as interleukin 8 (IL-8) (*Am. J. Respir. Crit. Care Med.* 156 1997 266-271) or interleukin 5 (IL-5) (EP 0775489 and EP 0771564) is known as well.

Macrolides have been found useful in the treatment of asthma, either because of their antimicrobial activity, or through a steroid-sparing effect. However, other properties such as eosinophil or neutrophil activation inhibition, have been suggested to explain the reduction in airway hyperresponsiveness. Respiratory and other inflammatory diseases (e.g. skin diseases or sinusitis) may also benefit from macrolide therapy (Labro M.T. *J. Antimicrob. Chemother.* 1998, 41,37; Cazzola m et al., *Mondaldi Arch Chest Dis* 200,55,231; Avila P.C. et al., *Ann Allergy Asthma Immunol* 200, 84 565; Amayasu H. Et al., *Ann Allergy Asthma Immunol* 2000, 84, 594; and Shoji T. Et al., *Clin Exp Allergy* 1999,29, 950; incorporated by reference in their entirety)

BRIEF DESCRIPTION OF THE FIGURES

Figure 1 represents efficacy of compound 38 and 40 in inducing hybridoma 13 apoptosis, as compared to standard dexamethasone.

Figure 2 and 3 represent inhibition of interleukin 4 and 5 respectively, by compound 31, as compared to standard dexamethasone.

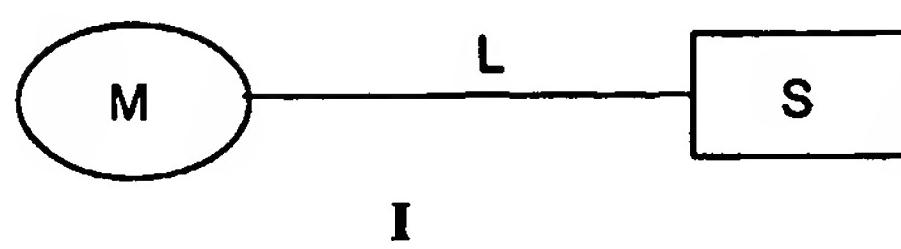
Figure 4 represents reduction in eosinophil percentage in BALF by compound 31, as compared with standard fluticasone propionate. Compound 31 was given i.n. in dose of 2 mg/kg, while fluticasone propionate was given 1 mg/kg.

TECHNICAL SOLUTION

Compounds represented by Formula I differ from steroids that are either unconjugated or conjugated to other types of molecules in that they combine the anti-inflammatory and immuno suppressant properties of steroids with the accumulation property of macrolides which are known to accumulate in cells effecting the inflammatory immune response that needs to be subdued. Such action of the compounds of Formula I arises from the macrolide moiety M which has the pharmacokinetic property of accumulating in immune system cells, notably phagocytes. This enables the compounds of Formula I to act predominantly if not exclusively at the inflammation site, by "riding" the macrolide within the very inflammation cells recruited to the locus of inflammation, and by inhibiting the production of inflammation mediators. In such a manner, the unfavourable systemic side-effects of corticosteroids are avoided or reduced and the activity of the compounds of the invention focused to the afflicted organ or tissue. After topical or systemic application, the hybrid molecules of the invention rapidly accumulate in inflammation cells, wherein they act by inhibiting the production of cytokines and chemokines as well as of other inflammation mediators, thus suppressing or inhibiting the inflammation.

The compounds of Formula I, which are the object of the present invention, their pharmacologically acceptable salts, solvates and prodrugs, and pharmaceutical compositions comprising them, have hitherto not been described, except for certain specific constructs comprising the macrolide FK-506 and the steroid dexamethasone, or the macrolide cyclosporin and the steroid dexamethasone or the macrolide pateamine A and the steroid dexamethasone. See WO 97/41255; Romo, D., *et al.*, *J. Am. Chem. Soc.*, 1998, 120:12237-12254; Griffith, E.C., *et al.* *Methods in Enzymology*, 2000, 328:89-110. However, these documents do not disclose use of any such compounds either as anti-inflammatory substances or as immunosuppressants of unwanted immune responses or as inhibitors of eosinophilic accumulation in inflammation tissues, nor do they describe compositions containing such compounds as therapeutics for administration to a mammal, including a human.

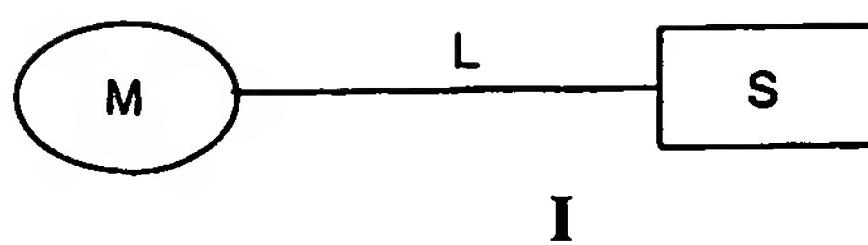
We have recently found that certain compounds within Formula I:



exert an improved therapeutic effect in the treatment of inflammation diseases, disorders and conditions. The symbol **M** in the above structure represents a macrolide subunit possessing the property of accumulation in inflammatory cells, **S** represents an anti-inflammatory steroid subunit and **L** represents a linker covalently linking **M** and **S**. Our co-pending commonly owned International Patent application PCT/HR02/00001 (incorporated herein by reference in its entirety, with a copy attached as Appendix A) describes compounds with the steroid subunit **S** linked via the chain **L** to position N/9a of 9-dihydro-9-deoxo-9a-aza-9a-homoerythromycin or to position C/3 of a des-cladinosyl azithromycin derivative or to position C/2' of the desozaminosyl group. However, hybrid compounds with the steroid subunit being linked with the position C/11 or N/9a with a modified or eliminated dimethylamino group on desozamine, which also possess the above mentioned therapeutic action, have not so far been described. Nor have hybrid compounds been described with the macrolide being linked to the steroid through position C/6 on the macrolide lactonic ring or through position C/4" of the cladinosyl group or through the amine group of position C/3' of the desozamine group or N/9a with linking to the steroid through 17 α -OH group of the steroid subunit. Moreover, in the foregoing PCT/HR02/00001 other types of steroid-macrolide hybrid compounds are not specifically described. All such compounds are the subject of the present application.

The present invention is directed to

- (a) new "hybrid" compounds represented by Formula I:



wherein **M** represents a macrolide subunit possessing the property of accumulation in inflammatory cells, **S** represents a steroid, as defined below, and **L** represents a linking group covalently linking **M** and **S**;

(b) compositions containing one or more of the foregoing compounds in an amount effective to combat inflammation and thereby to treat disorders and conditions involving inflammation in mammals, including humans; and

(c) methods for using these compounds to treat such disorders and conditions.

The present compounds advantageously provide an improved therapeutic effect and/or an improved side effect profile.

Suitable macrolide subunits for the hybrid compounds of the present invention can be selected without limitation from multi-member lactonic ring molecules, wherein "member" refers to the carbon atoms or heteroatoms in the ring, and "multi" is a number greater than about 10, preferably from 10 to about 50, more preferably 12-, 14-, 15-, 16-, 17- and 18-member lactonic ring macrolides. 14- and 15-member ring macrolide subunits are particularly preferred, with azithromycin and its derivatives and erythromycin and its derivatives being most preferred.

More specific nonlimiting examples of molecules from which the macrolide subunit can be selected are the following:

(i) Macrolide antibiotics, including azalides, for example erythromycin, dirithromycin, azithromycin, 9-dihydro-9-deoxy-9a-aza-9a-homoerythromycin, HMR 3004, HMR 3647, HMR 3787, josamycin, erythromycylamine, ABT 773, flurithromycin, clarithromycin, tylosin, tilmicosin, oleandomycin, desmycosin, CP-

163505, roxithromycin, miocamycin and rokitamycin and derivatives thereof, such as ketolides (e.g., 3-ketone), lactams (e.g., 8a- or 9a- lactams) and derivatives lacking one or more sugar moieties.

(ii) Macrolide immunosuppressants, such as FK 506, cyclosporin, amphotericin and rapamycin;

(iii) Macrolide antifungals with host cell inhibitory properties, such as bafilomycins, concanamycin, nystatin, natamycin, candicidin, filipin, etruscomycin, trichomycin.

Methodologies for the synthesis of the above macrolides not commercially available and synthetic manipulation of macrolides in general are known to those of ordinary skill in the art, or may be found in: Denis A. et al. Bioorg. & Med. Chem. Lett 1999, 9, 3075-3080; Agouridas C. et al. J. Med. Chem. 1998, 41, 4080-4100; and EP-00680967 (1998); Sun Or Y. et al. J. Med. Chem. 2000, 43, 1045-1049; US Patent-5747467 (1998); McFarland J. W. et al. J. Med. Chem. 1997, 40, 1041-1045; Denis A. et al. Bioorg.& Med. Chem. Lett. 1998, 8, 2427-2432 ;WO-09951616 (1999); Lartey et al. J Med Chem. 1995, 38, 1793-1798; EP 0984019; WO 98/56801, herein incorporated by reference in their entirety.

Additional suitable macrolides are known, some being disclosed in Bryskier, A. J., et al. *Macrolides, Chemistry, Pharmacology and Clinical Use*; Bryskier, Arnette Blackwell: Paris, 1993; pp 485-491, incorporated by reference in its entirety; in Ma, Z. et al. *Current Medicinal Chemistry - Anti-Infective Agents* 2002, 1, 15-34, also incorporated by reference in its entirety; in Romo, D. et al. *J. Am. Chem. Soc.* 1998, 120; 12237-12254; also incorporated by reference in its entirety. See, in particular the structures and derivatives for 14- and 16-member ring macrolides at pp 487-491 of Bryskier et al. and the various ketolide derivatives and syntheses in Ma et al. notably in all the structure tables and all the reaction schemes. All these macrolides after being conjugated to steroids are within the scope of the present invention. The foregoing specifically named macrolides as well as the ones referenced above are either commercially available or their structures and methods for their syntheses are known.

The structure and synthesis of the foregoing specifically enumerated compounds and several derivatives thereof is either known or well within the skill of the art.

In one subset of compounds according to the invention, if one constituent of the present compounds is FK-506 or cyclosporine, the other cannot be dexamethasone. In a further subset of the present compounds, FK-506, cyclosporine and pateamine A are excluded from the macrolide subunits altogether. However these provisos do not extend necessarily to novel methods for using the compounds defined above (including those which have FK-506, cyclosporin or pateamine A as a constituent), or to pharmaceutical compositions and dosage forms containing effective amounts of the present compounds for administration to mammalian animals or humans, particularly against inflammation.

It is important that the macrolide subunit derive from a macrolide having the property of accumulating within immune system cells recruited to the site of inflammation, especially phagocytic cells. Additional examples of macrolides accumulating within specific classes of cells may be found in: Pascual A. et al. Clin. Microbiol. Infect. 2001, 7, 65-69. (Uptake and intracellular activity of ketolide HMR 3647 in human phagocytic and non-phagocytic cells). Hand W. L. et al. Int. J. Antimicrob. Agents, 2001, 18, 419-425. (Characteristics and mechanisms of azithromycin accumulation and efflux in human polymorphonuclear leukocytes); Amsden G. W. Int. J. Antimicrob. Agents, 2001, 18, 11-15. (Advanced-generation macrolides: tissue-directed antibiotics); Johnson J. D. et al. J. Lab. Clin. Med. 1980, 95, 429-439. (Antibiotic uptake by alveolar macrophages); Wildfeuer A. et al. Antimicrob. Agents Chemother. 1996, 40, 75-79. (Uptake of azithromycin by various cells and its intracellular activity under in vivo conditions); Scorneaux B. et al. Poult. Sci. 1998, 77, 1510-1521. (Intracellular accumulation, subcellular distribution, and efflux of tilmicosin in chicken phagocytes); Mtairag E. M. et al. J. Antimicrob. Chemother. 1994, 33, 523-536. (Investigation of dirithromycin and erythromycylamine uptake by human neutrophils in vitro); Anderson R. et al. J. Antimicrob. Chemother. 1988, 22, 923-933. (An in-vitro evaluation of the cellular uptake and intraphagocytic bioactivity of clarithromycin (A-56268, TE-031), a new

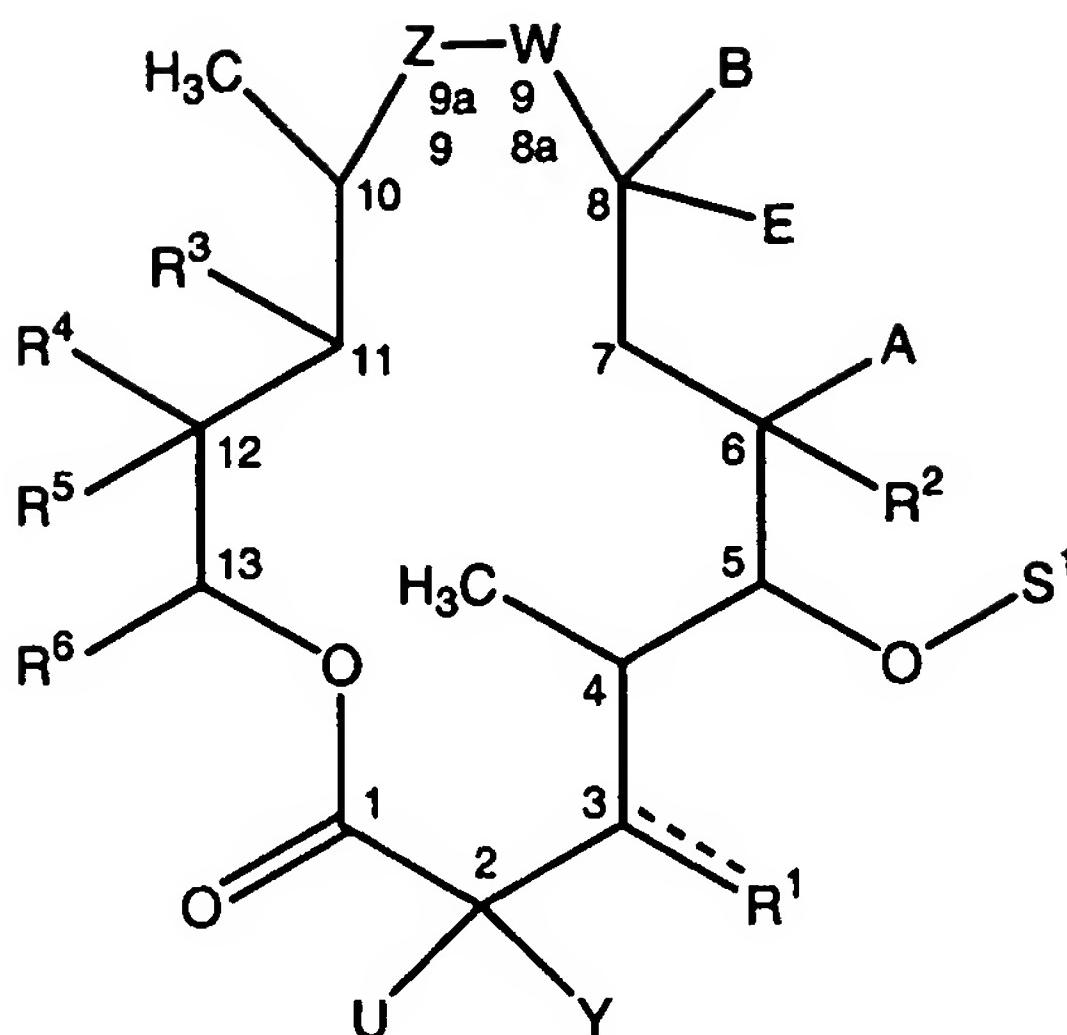
macrolide antimicrobial agent); Tasaka Y. et al. *Jpn. J. Antibiot.* 1988, 41, 836-840. (Rokitamycin uptake by alveolar macrophages); Harf R. et al. *J. Antimicrob. Chemother.* 1988, 22, 135-140. (Spiramycin uptake by alveolar macrophages) herein incorporated by reference in their entirety. Most of the lactonic compounds defined above are known to have this property, but if not, it can be easily tested by a person of ordinary skill in the field of the invention, using one of the well-known assays for this purpose. For example, the procedure detailed by Olsen, K. M. et al. *Antimicrob. Agents & Chemother.* 1996, 40, 2582-2585, which is hereby incorporated by reference. Briefly, the cells to be tested, e.g., polymorphonuclear leukocytes can be obtained from venous blood of healthy volunteers by Ficoll-Hypaque centrifugation followed by 2% dextran sedimentation. Erythrocytes are removed by osmotic lysis, and PMN are evaluated by Trypan blue exclusion. Alternatively, other cell fractions can be separated and similarly tested.

Tritiated macrolide compounds (e.g., 10 µM) are incubated with 2.5×10^6 cells for 120 minutes (37 °C, 5% CO₂, 90% relative humidity) and the cells are subsequently removed from compound-containing supernatant by centrifugation e.g., through a silicon oil-paraffin layer (86 vol. %: 14 vol. %). The amount of compound is determined, e.g., by scintillation counting, and a score significantly elevated above background indicates accumulation of the macrolide in the cells being tested. Alternatively, the compound is not radiolabelled but the amount of compound can be determined by HPLC.

Other assay methods that can be used are disclosed in Bryskier, A. J. et al. *Macrolides, Chemistry, Pharmacology and Clinical Use*; Arnette Blackwell: Paris, 1993; pp 375-386, incorporated by reference. See, in particular phagocytic uptake determination at pp 380-381 and the particular descriptions as to uptake and localization of macrolides at pp 381, 383 and 385 and the tables at 382.

In some preferred embodiments, this invention relates to compounds, their salts and solvates represented by Formula I, wherein M specifically represents a 14- or 15- member lactonic ring macrolide subunit more preferably represented by Formula II:

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wherein

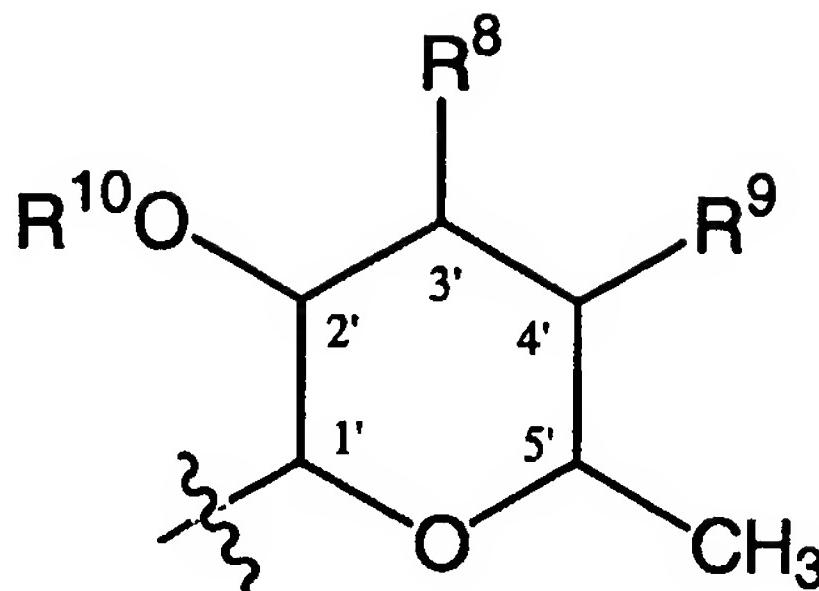
(i) Z and W independently are $\text{C}=\text{O}$, CH_2 , $\text{CH}-\text{NR}_t\text{R}_s$, NR_N , $\text{C}=\text{NR}_M$, or a bond, wherein

R_t and R_s independently are H or alkyl (preferably methyl or H);
 R_M is OH, OR^P, alkoxy or substituted alkoxy (in either Syn or Anti configurations or mixtures thereof)
 R_N is H, R^P, alkyl, alkenyl, alkynyl, alkoxy, alkoxyalkyl, or -C(=X)-NR_tR_s; and
X is O or S;

provided that Z and W cannot both simultaneously be $\text{C}=\text{O}$, $\text{CH}-\text{NR}_t\text{R}_s$,

CH_2 , NR_N , $\text{C}=\text{NR}_M$, or a bond,

- (ii) U and Y are independently H, halogen, alkyl, or hydroxymethyl (preferably H, methyl, or hydroxymethyl);
(iii) R¹ is hydroxy, OR^P, -O-S², or =O;
(iv) S¹ is a sugar moiety at position C/5 (e.g., a desozamine group) of the formula:



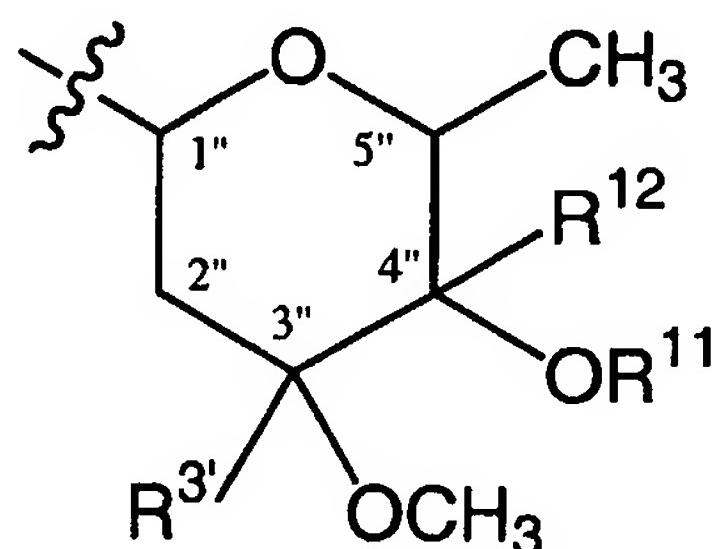
wherein

R^8 and R^9 are both hydrogen or together form a bond, or R^9 is hydrogen and R^8 is $-N(CH_3)R^y$, wherein

R^y is R^p , R^z or $-C(O)R^z$, wherein R^z is hydrogen or cycloalkyl (preferably cyclohexyl) or alkyl (preferably a C₁-C₇ alkyl) or alkenyl (preferably C₂-C₇-alkenyl) or alkynyl (preferably C₂-C₇-alkynyl) aryl or heteroaryl or alkyl substituted with C₂-C₇ alkyl, C₂-C₇ alkenyl, C₂-C₇ alkynyl, aryl or heteroaryl (R^y is preferably hydrogen, methyl, ethyl, n-propyl, i-propyl, n-butyl, $-C(O)CH_3$, $-CH_2$ -phenyl, or cyclohexyl);

R^{10} is hydrogen or R^p ;

(v) S² sugar moiety (e.g., is a cladinosyl group) of the formula



wherein $R^{3'}$ is H or methyl and R^{11} is hydrogen or R^p or $O-R^{11}$ is a group that with R^{12} and with C/4'' carbon atom forms a $>C=O$ or epoxy group; R^{12} is hydrogen or a group that with $O-R^{11}$ and with C/4'' carbon atom forms a $>C=O$ or epoxy group;

(vi) R^2 is H, hydroxy, OR^p group, alkoxy (preferably C₁-C₄ alkoxy, most preferably methoxy) or substituted alkoxy;

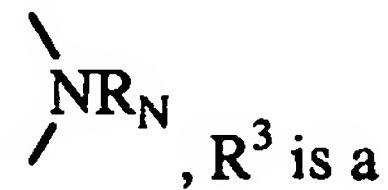
(vii) A is H or methyl;

(viii) B is methyl or epoxy;

(ix) E is H or halogen (preferably fluorine);

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(x) R^3 is hydroxy, OR^P group or alkoxy (preferably C_1-C_4 alkoxy, most preferably methoxy), substituted alkoxy or R^3 is a group that can combine with R^5 to

form a "bridge" (e.g., a cyclic carbonate or carbamate) or if W or Z is , R^3 is a group that can combine with W or Z to form a "bridge" (e.g., a cyclic carbamate);

(xi) R^4 is C_1-C_4 alkyl (preferably methyl);

(xii) R^5 is H, hydroxy, OR^P group, C_1-C_4 alkoxy, substituted alkoxy or a group that may combine with R^3 to form a bridge (e.g., a cyclic carbonate or carbamate);

(xiii) R^6 is H or C_1-C_4 alkyl (preferably methyl or ethyl);

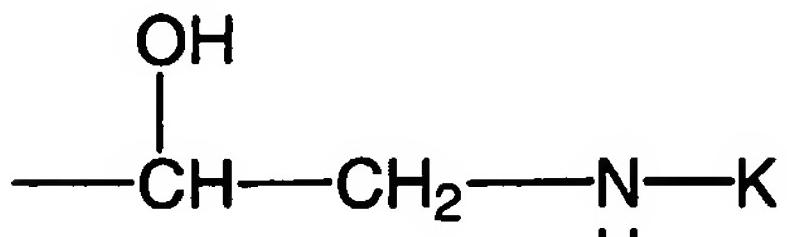
wherein the subunit M has a linkage site through which it is linked to the subunit S via the linking group L, the linkage site being at one or more of the following:

a. any reactive hydroxy, N, or epoxy group located on S^1 , S^2 , or an aglycono oxygen if S^1 or S^2 is cleaved off;

b. a reactive $>N-R_N$ or $-NR_tR_s$ or $=O$ group located on Z or W;

c. a reactive hydroxy group located at any one of R^1 , R^2 , R^3 , and R^5 ;

d. any other group that can be first derivatized to a hydroxy or $-NR_tR_s$ group and then linked to K (e.g., $OH \rightarrow =O \rightarrow$ epoxy \rightarrow



) wherein K is the part of the linking molecule L.

One or more R^P groups may be independently present in the macrolide subunit of Formula II, wherein R^P represents a protective group which may be selected from alkyl (preferably methyl), alkanoyl (preferably acetyl), alkoxycarbonyl (preferably methoxycarbonyl or *tert*-butoxycarbonyl), arylmethoxycarbonyl (preferably benzyloxycarbonyl), aroyl (preferably benzoyl), arylalkyl (preferably benzyl), alkylsilyl (preferably trimethylsilyl) or alkylsilylalkoxyalkyl (preferably trimethylsilylethoxymethyl). The amino protecting groups may be removed by conventional techniques. Thus, for example acyl groups like alkanoyl, alkoxycarbonyl or aroyl may be removed by solvolysis, e.g. by hydrolysis under acidic or basic

conditions. Arylmethoxycarbonyl group (benzyloxycarbonyl) may be cleaved by hydrogenolysis in the presence of a catalyst such as palladium-on-charcoal.

L represents the structure **VA** or **VB**:

VA	$X^1-(CH_2)_m-X^2$ or
VB	$X^1-(CH_2)_m-Q-(CH_2)_n-X^2$

wherein

X^1 is selected from: -CH₂, -CH₂-NH-, -C(O)-, -OC(O)-, =N-O-, -OC(O)NH- or -OC(O)NH-;

X^2 is selected from: -NH-, -CH₂-; -NHC(O)-, -C(=O) or -OC(O)-;

Q is -NH- or -CH₂-;

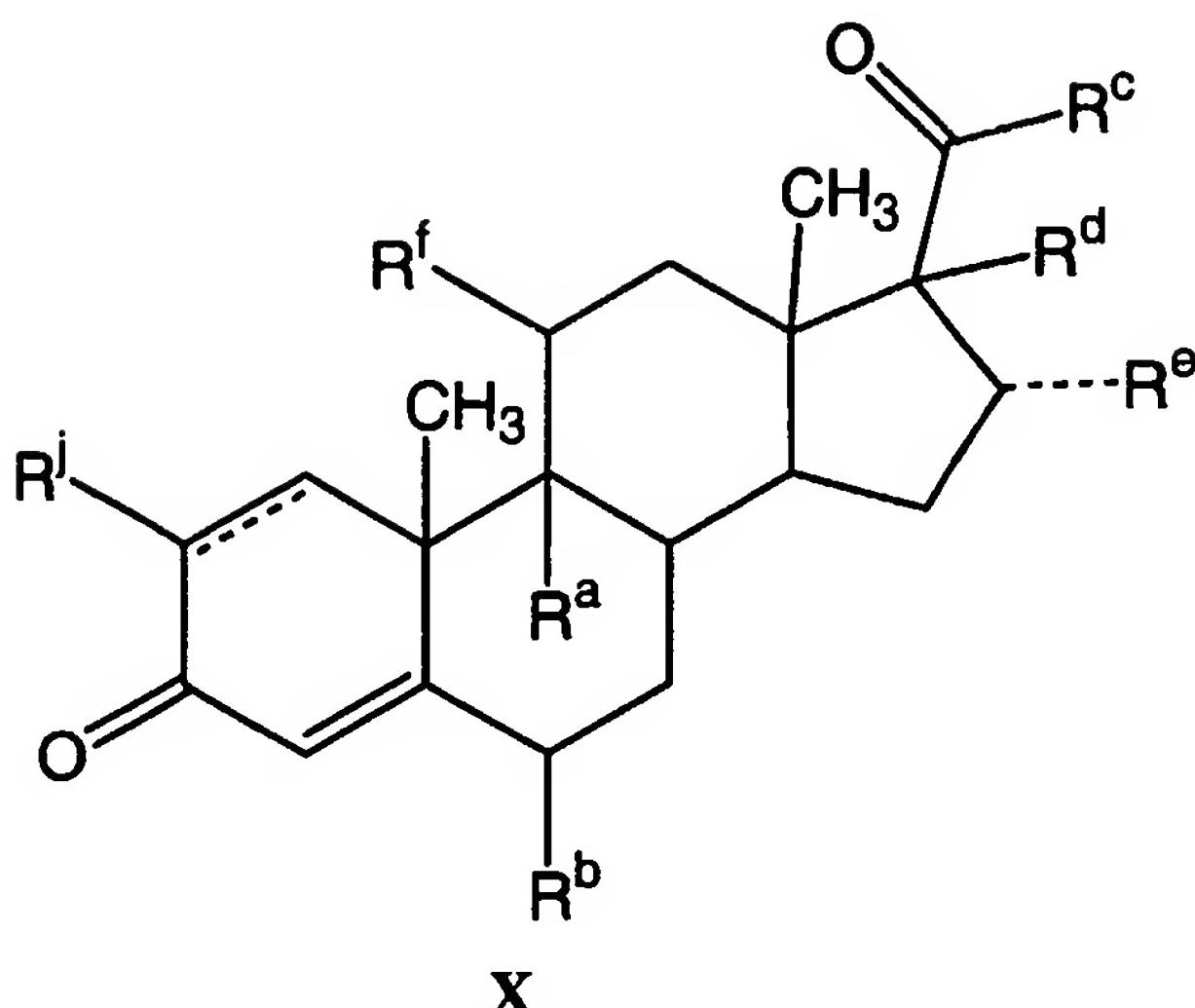
wherein each -CH₂- or -NH- group are optionally substituted by C₁-C₇-alkyl, C₂-C₇-alkenyl, C₂-C₇-alkynyl, C(O)R^x, C(O)OR^x, C(O)NHR^x wherein R^x may be C₁-C₇-alkyl, aryl or heteroaryl;

the symbols m and n are independently a whole number from 0 to 8

with the proviso that if Q=NH; n cannot be zero.

This definition of the linking group is preferred not only for hybrids of steroids and macrolides of Formula II but for any conjugate within Formula I. Other linking groups can be used as long as they provide the necessary spacer and can serve to link one subunit of the Formula I with the other, as is well-known in the art. See, e.g., U.S. Patent 6,297,260, which is incorporated by reference in its entirety, especially its claim 1 and the specific list of steroids.

S is a steroid subunit, preferably of Formula X:



wherein

R^a and R^b are, independently of each other, hydrogen or halogen;

R^c is hydroxy, alkoxy (preferably methoxy), substituted alkoxy, alkyl, thiocarbamoyl, carbamoyl or a valence-bond;

R^d and R^e are, independently of each other, hydrogen, OH, CH_3 or C_1-C_4 alkoxy (preferably methoxy or n-propoxy) or each are a group that forms a 1,3-dioxolane ring with the other (optionally alkyl or alkenyl mono-or di-substituted) (preferably a 2,2-dimethyl or 2-monopropyl or trans-propenyl ring) or a valence bond;

R^f is hydrogen, hydroxy, chloro, or $=O$ forming a keto group with the carbon atom it is attached to;

R^j is hydrogen or chloro

and their pharmacologically acceptable salts and solvates.

Alternatively with the present invention are steroid subunits disclosed in WO 94/14834, wherein instead of the group $>CH-C(O)-R^c$ they have the group $>CH-S(O)_n-R^c$ wherein n is an integer of 0 to 2. See WO 94/14834 incorporated in its entirety by reference, especially pp 2-3.

More generally, steroids useful as a source of steroid subunits include, but are not limited to, corticosteroids (such as glucocorticoids and mineralocorticoids) and androgens. Non-limiting examples of corticosteroids include cortisol, cortisone,

clobetasol, hydrocortisone, fludrocortisone, fludroxycortide, flumetasone, flunisolide, fluocinolone, fluocinonide, fluocortolone, fluorometholone, prednisone, prednisolone, 6-alpha-methylprednisolone, triamcinolone, alclometasone, beclometasone, betamethasone, budesonide, dexamethasone, amcinonide, cortivazol, desonide, desoximethasone diflucortolone, difluprednate, fluclorolone and dichlorisone, flupernidene, fluticasone, halcinonide, meprednisone, methylprednisolone, paramethasone, prednazoline, prednylidene, tixocortol, triamcinolone, and acid derivatives thereof, e.g., acetate, propionate, dipropionate, valerate, phosphate, isonicotinate, metasulfobenzoate, tebutate, and hemisuccinate).

The symbols M, L and S represent three different subunits of the compounds represented by the structure I. The symbol M represents the macrolide subunit and the symbol S the steroid subunit which is linked via the linker molecule L with a linkage site of the subunit M. As discussed above, when M is an azithromycin lactone ring then the linkage is preferably effected through position C/3 of the aglycone moiety or at position C/6, C/11 or at position N/9a of the compound of Formula II, or through position C/4" of the S² group or through the amine at 3' or 2' of the S¹ group.

Bold-faced bonds in the formulas presented herein denote bonds raised above the paper plane (β -position). Dash-drawn bonds denote bonds situated below the paper plane (α -position). Parallel full and broken lines denote bonds that can be either single or double. A single broken line denotes a bond either above or below the paper plane.

Unless stated otherwise, the following terms have the meanings ascribed to them below.

"Halogen" means a halogen atom which may preferably be: fluorine, chlorine or bromine (the most preferably fluorine or chlorine).

"Alkyl" means a linear or branched saturated monovalent hydrocarbon radical of one to ten carbon atoms, more preferably one to six carbon atoms. The preferred straight-chain or branched-chain alkyls include methyl, ethyl, propyl, iso-propyl, butyl, sec-butyl and tert-butyl. Methyl is most preferred. Alkyl groups may be

substituted with one up to five substituents including halogen (preferably fluorine or chlorine), hydroxy, alkoxy (preferably methoxy or ethoxy), acyl, acylamino cyano, amino, N-(C₁-C₄)alkylamino (preferably N-methylamino or N-ethylamino), N,N-di(C₁-C₄-alkyl)amino (preferably dimethylamino or diethylamino), aryl (preferably phenyl) or heteroaryl, thiocarbonylamino, acyloxy, amino, amidino, alkylamidino, thioamidino, aminoacyl, aminocarbonylamino, aminothiocarbonylamino, aminocarbonyloxy, aryl, heteroaryl, aryloxy, aryloxyaryl, nitro, carboxyl, carboxylalkyl, carboxyl-substituted alkyl, carboxyl-cycloalkyl, carboxyl-substituted cycloalkyl, carboxylaryl, carboxyl-substituted aryl, carboxylheteroaryl, carboxyl-substituted heteroaryl, carboxylheterocyclic, carboxyl-substituted heterocyclic, cycloalkyl, cycloalkoxy, heteroaryloxy, heterocyclyloxy, and oxycarbonylamino. Such substituted alkyl groups are within the present definition of "alkyl." The present definition of alkyl carries over to other groups having an alkyl moiety such as alkoxy.

"Alkenyl" means a linear or branched monovalent hydrocarbon radical of two to ten and preferably two to six carbon atoms which has at least one double carbon-carbon bond. Alkenyl groups may be substituted with the same groups as alkyl and such optionally substituted alkenyl groups are encompassed within the term "alkenyl". Ethenyl, propenyl, butenyl and cyclohexenyl are preferred.

"Alkynyl" means a linear or branched monovalent hydrocarbon radical, having a straight-chain or a branched-chain of two to ten, and preferably two to six carbon atoms and containing at least one and preferably no more than three triple carbon-carbon bonds. Alkynyl groups can be substituted with the same groups as alkyl, and the substituted groups are within the present definition of alkynyl. Ethynyl, propynyl and butynyl groups are preferred.

"Cycloalkyl" means a cyclic group having 3-8 carbon atoms having a single ring optionally fused to an aryl or heteroaryl group. The cycloalkyl groups can be substituted as specified for "aryl" below, and the substituted cycloalkyl groups are within the present definition of "cycloalkyl". Preferred cycloalkyls are cyclopentyl and cyclohexyl.

"Aryl" means an unsaturated aromatic carbocyclic group having 6-14 carbon atoms having a single ring such as phenyl or multiple fused rings such as naphthyl. Aryl may optionally be further fused to an aliphatic or aryl group or can be substituted with one or more substituents such as halogen (fluorine, chlorine and/or bromine), hydroxy, C₁-C₇ alkyl, C₁-C₇ alkoxy or aryloxy, C₁-C₇ alkylthio or arylthio, alkylsulfonyl, cyano or primary or nonprimary amino.

"Heteroaryl" means a monocyclic or a bicyclic aromatic hydrocarbon ring having from 2 to 10 carbon atoms and from 1 to 4 heteroatoms, such as O, S or N. The heteroaryl ring may optionally be fused to another heteroaryl, aryl or aliphatic cyclic group. Examples of this type are furan, thiophene, imidazole, indole, pyridine, oxazole, thiazole, pyrrole, pyrazole, tetrazole, pyrimidine, pyrazine and triazine, with furan, pyrrole, pyridine and indole being preferred. The term includes groups that are substituted with the same substituents as specified for aryl above.

"Heterocyclic" means a saturated or unsaturated group having a single or multiple rings and from 1 to 10 carbon atoms and from 1-4 heteroatoms selected from nitrogen, sulfur or oxygen, wherein in a fused ring system the other ring or rings can be aryl or heteroaryl. Heterocyclic groups can be substituted as specified for alkyl groups and the thus substituted heterocyclic groups are within the present definition.

When R^c represents a valence-bond, the steroid subunit S is linked via R^c with the chain L to the macrolide subunit M. The symbol K is sometimes used to refer to the part of the L group, linked to M, or to S at the context requires.

In the preparation of the compounds represented by the structure I, of the specified pharmacological activity, certain new compounds were prepared as intermediates in the preparation of pharmacologically active compounds. The present invention also relates to such intermediates.

The present invention also encompasses pharmaceutically acceptable salts of the present compounds. Pharmaceutically suitable salts of the compounds of the present invention include salts with inorganic acids (e.g. hydrochloric, hydrobromic, phosphoric, metaphosphoric, nitric or sulfuric acid) or organic acids (e.g. tartaric,

acetic, mesylic, trifluoroacetic, citric, maleic, lactic, fumaric, benzoic, succinic, methanesulfonic, oxalic and p-toluenesulfonic acids).

A further aspect of the present invention are the solvates (preferably hydrates) formed by the compounds represented by the structure I or their salts.

The present invention also encompasses pharmaceutically acceptable salts of the present compounds. Pharmaceutically suitable salts of the compounds of the present invention include salts with inorganic acids (hydrochloric, hydrobromic, phosphoric, metaphosphoric, nitric or sulfuric acid) or organic acids (tartaric, acetic, methane-sulfonic, trifluoroacetic, citric, maleic, lactic, fumaric, benzoic, succinic, methanesulfonic, oxalic and p-toluenesulfonic acids).

The present invention also encompasses prodrugs of Formula I compounds, i.e., compounds which release an active parent drug according to Formula I *in vivo* when administered to a mammalian subject. Prodrugs of a compound of Formula I are prepared by modifying functional groups present in the compound of Formula I in such a way that the modifications may be cleaved *in vivo* to release the parent compound. Prodrugs include compounds of Formula I wherein a hydroxy, amino, or carboxy group of a Formula I compound is bonded to any group that may be cleaved *in vivo* to regenerate the free hydroxyl, amino or carboxy group, respectively. Examples of prodrugs include, but are not limited to esters (e.g., acetate, formate, and benzoate derivatives) of compounds of Formula I.

The compounds of Formula I have one or more chirality centers and, depending on the nature of individual substituents, they can also have geometrical isomers. Isomers that differ in the arrangement of their atoms in space are termed "stereoisomers". Stereoisomers that are not mirror images of one another are termed "diastereomers" and those that are non-superimposable mirror images of each other are termed "enantiomers". When a compound has a chiral center, a pair of enantiomers is possible. An enantiomer can be characterized by the absolute configuration of its asymmetric center and is described by the R- and S-sequencing rules of Cahn and Prelog, or by the manner in which the molecule rotates the plane of polarized light and designated as dextrorotatory or levorotatory (i.e., as (+) or (-))

isomer respectively). A chiral compound can exist as either an individual enantiomer or as a mixture of enantiomers. A mixture containing equal proportions of the enantiomers is called a "racemic mixture". The present invention encompasses all individual isomers of compounds of Formula I. The description or naming of a particular compound in the specification and claims is intended to include both individual enantiomers and mixtures, racemic or otherwise, thereof. Methods for the determination of stereochemistry and the separation of stereoisomers are well-known in the art.

The present invention also encompasses stereoisomers of the syn-anti type, encountered when an oxime or similar group is present. The group of highest Cahn Ingold Prelog priority attached to one of the terminal doubly bonded atoms of the oxime, is compared with hydroxyl group of the oxime. The stereoisomer is designated as *Z* (*zusammen* = together) or *Syn* if the oxime hydroxyl lies on the same side of a reference plane passing through the C=N double bond as the group of highest priority; the other stereoisomer is designated as *E* (*entgegen* = opposite) or *Anti*.

A "pharmaceutically acceptable excipient" means an excipient that is useful in preparing a pharmaceutical composition that is generally safe, non-toxic and neither biologically nor otherwise undesirable, and includes an excipient that is acceptable for veterinary use as well as human pharmaceutical use. A "pharmaceutically acceptable excipient" as used in the present application includes both one and more than one such excipient.

"Treating" or "treatment" of a state, disorder or condition includes:

- (1) preventing or delaying the appearance of clinical symptoms of the state, disorder or condition developing in a mammal that may be afflicted with or predisposed to the state, disorder or condition but does not yet experience or display clinical or subclinical symptoms of the state, disorder or condition,
- (2) inhibiting the state, disorder or condition, i.e., arresting or reducing the development of the disease or at least one clinical or subclinical symptom thereof, or

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(3) relieving the disease, i.e., causing regression of the state, disorder or condition or at least one of its clinical or subclinical symptoms.

The benefit to a subject to be treated is either statically significant or at least perceptible to the patient or to the physician

A "therapeutically effective amount" means the amount of a compound that, when administered to a mammal for treating a state, disorder or condition, is sufficient to effect such treatment. The "therapeutically effective amount" will vary depending on the compound, the disease and its severity and the age, weight, physical condition and responsiveness of the mammal to be treated.

The four classic symptoms of acute inflammation are redness, elevated temperature. Swelling, and pain in the affected area, and loss of function of the affected organ.

Symptoms and signs of inflammation associated with specific conditions include:

- rheumatoid arthritis- pain, swelling, warmth and tenderness of the involved joints; generalized and morning stiffness;
- insulin-dependent diabetes mellitus- insulitis; this condition can lead to a variety of complications with an inflammatory component, including: retinopathy, neuropathy, nephropathy; coronary artery disease, peripheral vascular disease, and cerebrovascular disease;
- autoimmune thyroiditis- weakness, constipation, shortness of breath, puffiness of the face, hands and feet, peripheral edema, bradycardia;
- multiple sclerosis- spasticity, blurry vision, vertigo, limb weakness, paresthesias;
- uveoretinitis- decreased night vision, loss of peripheral vision;
- lupus erythematosus- joint pain, rash, photosensitivity, fever, muscle pain, puffiness of the hands and feet, abnormal urinalysis (hematuria, cylindruria, proteinuria), glomerulonephritis, cognitive dysfunction, vessel thrombosis, pericarditis;

- scleroderma- Raynaud's disease; swelling of the hands, arms, legs and face; skin thickening; pain, swelling and stiffness of the fingers and knees, gastrointestinal dysfunction, restrictive lung disease; pericarditis,; renal failure;
- other arthritic conditions having an inflammatory component such as rheumatoid spondylitis, osteoarthritis, septic arthritis and polyarthritis- fever, pain, swelling, tenderness;
- other inflammatory brain disorders, such as meningitis, Alzheimer's disease, AIDS dementia encephalitis- photophobia, cognitive dysfunction, memory loss;
- other inflammatory eye inflammations, such as retinitis- decreased visual acuity;
- inflammatory skin disorders, such as , eczema, other dermatites (e.g., atopic, contact), psoriasis, burns induced by UV radiation (sun rays and similar UV sources)- erythema, pain, scaling, swelling, tenderness;
- inflammatory bowel disease, such as Crohn's disease, ulcerative colitis- pain, diarrhea, constipation, rectal bleeding, fever, arthritis;
- asthma- shortness of breath, wheezing;
- other allergy disorders, such as allergic rhinitis- sneezing, itching, runny nose
- conditions associated with acute trauma such as cerebral injury following stroke- sensory loss, motor loss, cognitive loss;
- heart tissue injury due to myocardial ischemia- pain, shortness of breath;
- lung injury such as that which occurs in adult respiratory distress syndrome- shortness of breath, hyperventilation, decreased oxygenation, pulmonary infiltrates;
- inflammation accompanying infection, such as sepsis, septic shock, toxic shock syndrome- fever, respiratory failure, tachycardia, hypotension, leukocytosis;
- other inflammatory conditions associated with particular organs or tissues, such as nephritis (e.g., glomerulonephritis)-oliguria, abnormal urinalysis; inflamed appendix- fever, pain, tenderness, leukocytosis;gout- pain, tenderness, swelling and erythema of the involved joint, elevated serum and/or

urinary uric acid; inflamed gall bladder- abdominal pain and tenderness, fever, nausea, leukocytosis; chronic obstructive pulmonary disease- shortness of breath, wheezing; congestive heart failure- shortness of breath, rales, peripheral edema; Type II diabetes- end organ complications including cardiovascular, ocular, renal, and peripheral vascular disease lung fibrosis- hyperventilation, shortness of breath, decreased oxygenation; vascular disease, such as atherosclerosis and restenosis- pain, loss of sensation, diminished pulses, loss of function and alloimmunity leading to transplant rejection- pain, tenderness, fever.

Subclinical symptoms include without limitation diagnostic markers for inflammation the appearance of which may precede the manifestation of clinical symptoms. One class of subclinical symptoms is immunological symptoms, such as the invasion or accumulation in an organ or tissue of proinflammatory lymphoid cells or the presence locally or peripherally of activated pro-inflammatory lymphoid cells recognizing a pathogen or an antigen specific to the organ or tissue. Activation of lymphoid cells can be measured by techniques known in the art.

“Delivering” a therapeutically effective amount of an active ingredient to a particular location within a host means causing a therapeutically effective blood concentration of the active ingredient at the particular location. This can be accomplished, e.g., by local or by systemic administration of the active ingredient to the host.

The present invention also relates to possible tautomeric forms which can be formed by individual compounds of Formula I.

Further preferred are compounds of Formula I wherein **M** represents a group of Formula II wherein Z is >N-R_N wherein R_N is hydrogen or a methyl; W is >CH₂; B is methyl; E is hydrogen; R² is hydroxy; A is methyl; R⁸ at the C/3' position in S¹ is selected from: hydrogen, amino, N-methylamino, N,N-dimethylamino, N-methyl-N-

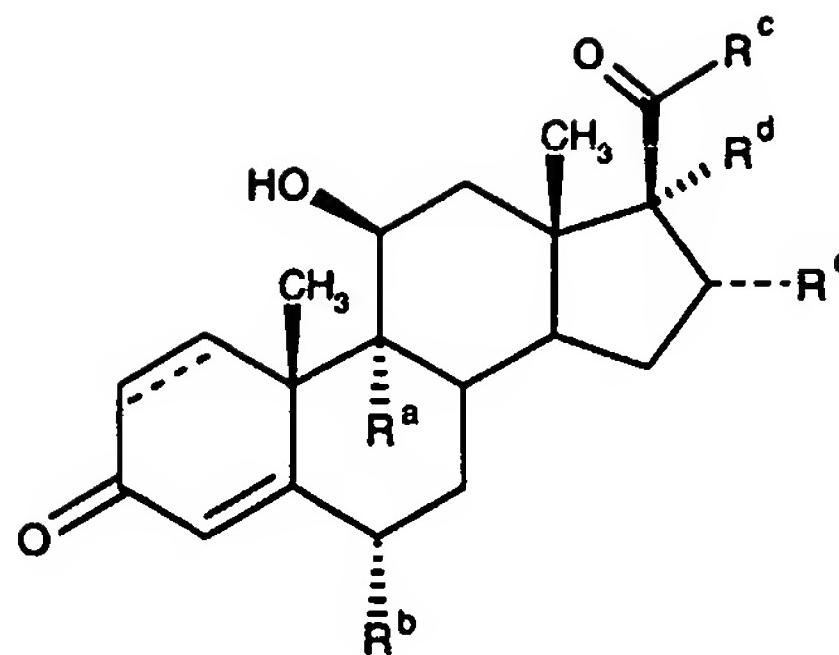
(C₂-C₄)-alkylamino, N-methyl-N-benzylamino or N-methyl-N-cyclohexyl-amino; U=H, Y=CH₃, R¹ is O-S² group wherein S² is a cladinosyl group wherein the 3" position is both methyl and methoxy substituted, R⁶ is ethyl, R⁵ is methyl; R¹¹ and R¹² are hydrogen and R¹³ is methyl; R⁴ is OH or forms a cyclic carbonate bridge with R³, R³ is OH or forms a cyclic carbonate bridge with R⁴, and the 3' position of the desozamine group is H or -N(CH₃)R^y as defined above. The linkage is through the N of Z, or through the O of R³. In the latter case, Z is NH.

Also preferred are compounds of Formula I wherein M represents a group of Formula II wherein: Z is >NH or >NCH₃ or >NC(O)NHR^x wherein R^x is iso-propyl; W is >C=O or >CH₂ (provided that when Z is >NCH₃ W cannot be >C=O); R² is hydroxy or methoxy; A is methyl; E is hydrogen; B is methyl; R³ is hydroxy; R⁵ is methyl; R⁴ is hydroxy or methoxy; R⁶ is ethyl; R¹ is a O-S² group wherein S² is a cladinosyl group wherein R¹³ is methyl; R¹¹ is hydrogen, or O-R¹¹ is a group that forms with R¹² and with C/4" carbon atom a >C=O or epoxy group, R¹² is hydrogen or a group that forms with O-R¹¹ and with a C/4" carbon atom a >C=O or epoxy group; C/3' position of the S¹ group is hydrogen, amino or N-(C₁-C₆)-alkylamino or N,N-(C₁-C₆)-dialkylamino. The linkage is through the amino of R⁸ at the C/3' position or through the oxygen of R² at C/6 position or through the carbon of R¹² or the oxygen of R¹¹ both at C/4" position.

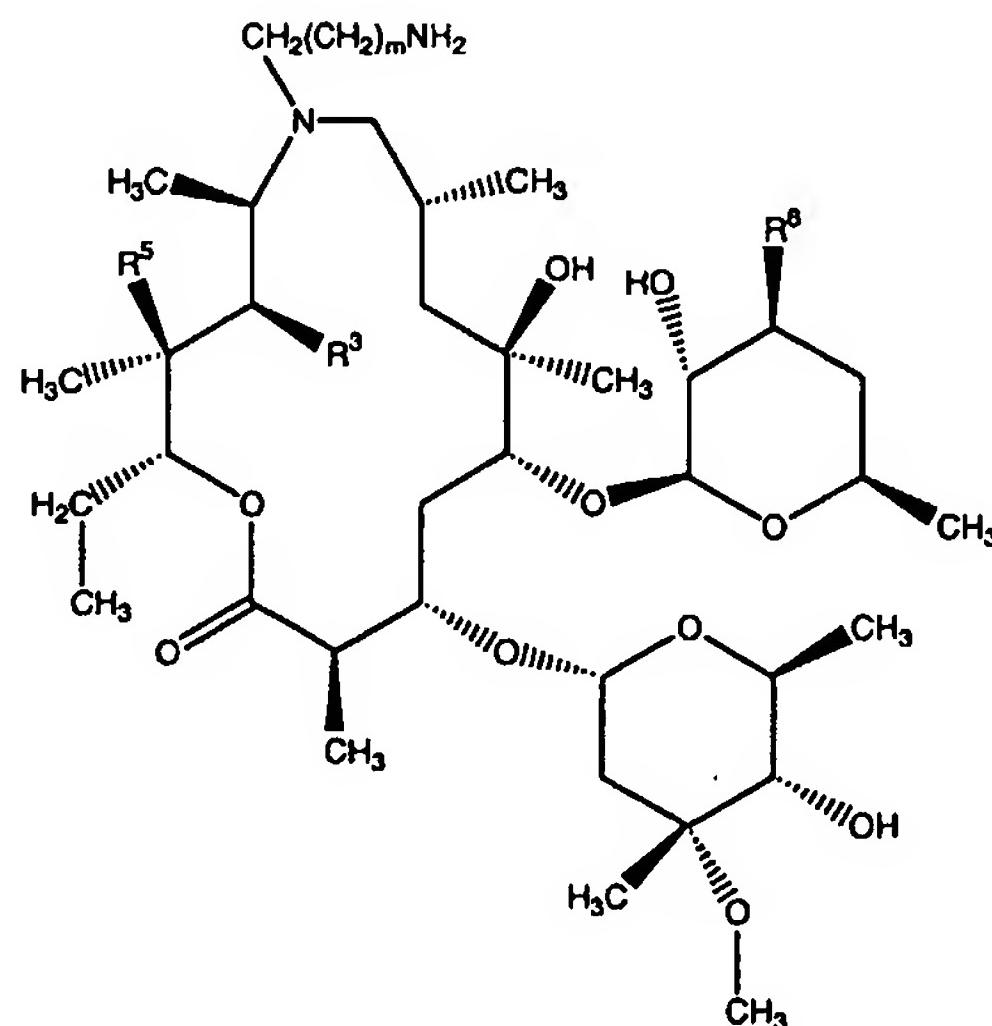
Methods of Preparation A

A further aspect of the present invention relates to a method for the preparation of specific compounds within Formula I comprising:

- a) for the compounds of Formula I, wherein L has X¹ being -CH₂-, X² being -NH-, and Q is absent (Formula VA), a reaction of carboxylic acid of the steroid subunit of Formula Xa:

**Xa**

wherein $-\text{C}(\text{O})\text{R}^{\text{c}}$ is an activated carboxylic acid and R^{a} , R^{b} , R^{d} , and R^{e} are defined as in Formula X, with a free amino group of the macrolide subunit of Formula VIa:

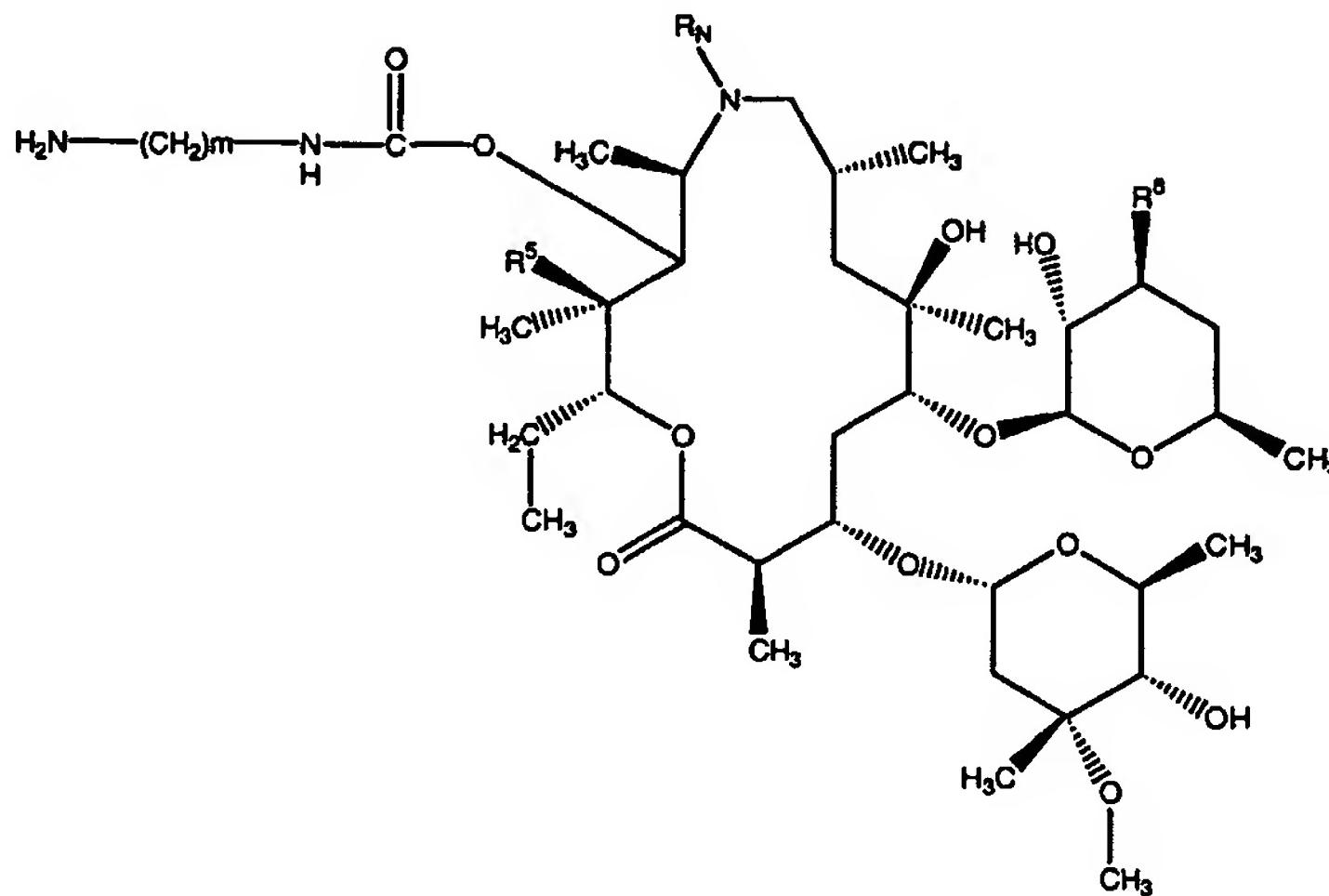
**VIa**

wherein R^{3} , R^{5} , and R^{8} are defined in connection with Formula II above;

b) for the compounds of Formula I, wherein X^{1} in L is $-\text{C}(\text{O})\text{NH}-$ and X^{2} is $-\text{NH}-$ (also Formula VA), the reaction of the carboxylic acid of the

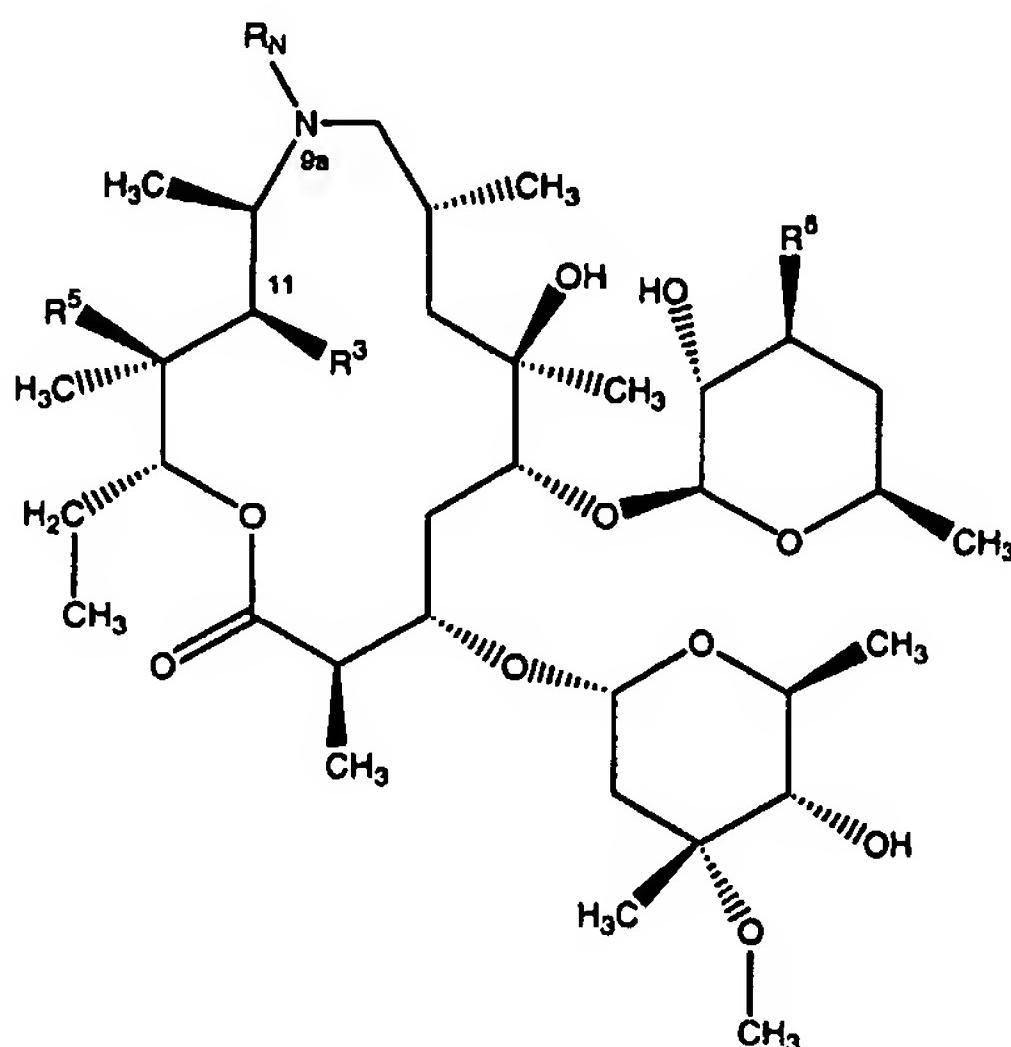
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steroid subunit of the structure **X**, wherein $-C(O)R^c$ is an activated carboxylic acid, with a free amino group of the macrolide subunit of the structure **VIb**:

**VIb**

wherein R_N , R^5 , and R^8 are as defined in Formula **II** and m is as defined in Formula **VA**.

c) for the compounds of Formula **I**, wherein R^8 is H, $-NH-CH_3$, or $-N(CH_3)R^y$ by removal or change of the $-N(CH_3)_2$ group on the desozamino sugar of a compound of Formula **I** having a macrolide subunit of Formula **VIc**:



wherein R_N, R³, and R⁸ are as defined in Formula II.

Methods of Preparation:

a) A compound within Formula I is prepared by a reaction of a carboxylic acid of the steroid subunit of the structure X and the amino group of the macrolide subunit of the structure VIa according to the process as described in International Application No PCT/HR02/00001, whereby the amide linkage is effected, using the usual acid derivatives having an activating effect on carboxylic acid, such as anhydrous halogenides with mixed anhydrides especially carbodiimides or benzotriazole (e.g. hydroxybenzotriazole). The reaction proceeds in the presence of a base (preferably an organic), e.g.. triethylamine, at room temperature under an inert atmosphere, e.g., argon blanket over the period from several hours to several days. (D. Romo et al., *J. Am. Chem. Soc.*, **1998**, *120*, 12237).

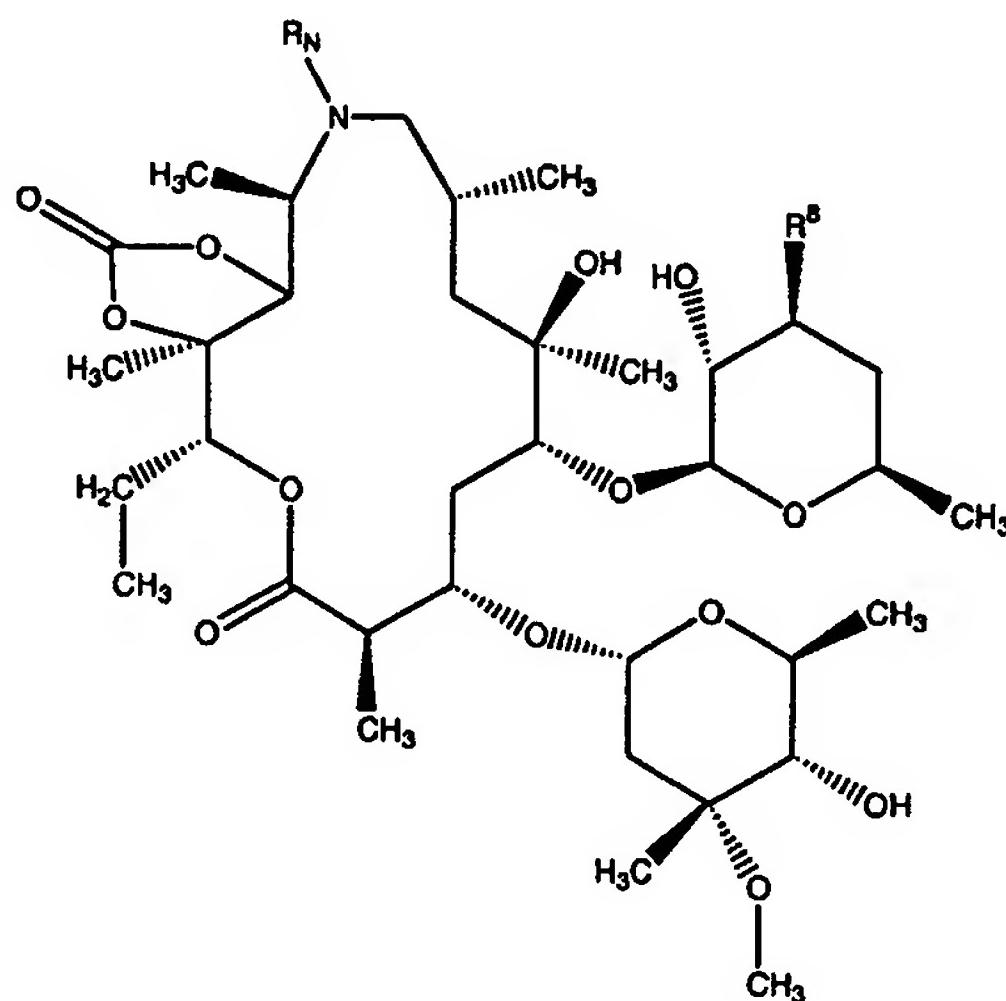
Steroid subunits of the structure X are either commercially available products or have been obtained by known methods (Suzuki, T. et al., *Chem. Soc., Perkin Trans. 1* **1998**, 3831-3836.), (McLean, H.M. et al., *J. Pharm. Sci.* **1994**, *83*, 476-480.), (Little, R.J. et al., *Pharm. Res.* **1999**, *16*, 961-967.), (Kertesz D.J. et al., *J. Org. Chem.* **1986**, *51*, 2315-2328.), (Bodor, N.S., *US Patent 4,710,495* **1987**), (Phillipps, G. et al., *J. Med. Chem.* **1994**, *37*, 3717-3729) all incorporated in their entirety by reference.

Starting macrolide subunits of the structure VIa may be obtained by the action of corresponding cyanoalkenyls (preferably acrylonitriles) on the corresponding macrolide subunits and then by hydrogenation of the formed nitrile, according to the procedures of Bright, U.S. Patent No. 4,474,768, October 2, 1984; and Bright, G.M. et al., 1998, *J. Antibiot.* **41**:1029-1047.

b) Preparation of a compound of Formula I is carried out by a reaction of the carboxylic acid of the steroid subunit represented by the structure X and the amino group of the macrolide subunit of the structure VIb, whereby the amine bond is

prepared, and the process proceeds in the manner and under conditions as described in the method a) above.

Starting macrolide subunits of the structure **VIIb** may be obtained by the action of the corresponding diaminoalkane (preferably diaminobutane) on the macrolide subunit **VII**:



VII

which can be obtained in by the action of ethylcarbonate on the macrolide subunit, as reported in the literature (EP 0984019 A1, which is hereby incorporated by reference).

c) A compound within Formula I is prepared by modification or elimination of the $-N(CH_3)_2$ group R^8 of the desozamine sugar of compounds having a macrolide subunit represented by Formula VIc. Modification can be effected by demethylation of one of the two methyl groups, e.g., by UV radiation. By alkylation or acylation of the obtained compound of Formula I, wherein R^8 is a $NH(CH_3)$ group, a new class of compounds of Formula I is formed, wherein R^8 is $-N(CH_3)R^2$. It is possible to eliminate the $-N(CH_3)_2$ group by oxidizing it and by forming of a corresponding N-oxide, which on heating yields the compounds of Formula I, wherein R^8 is hydrogen.

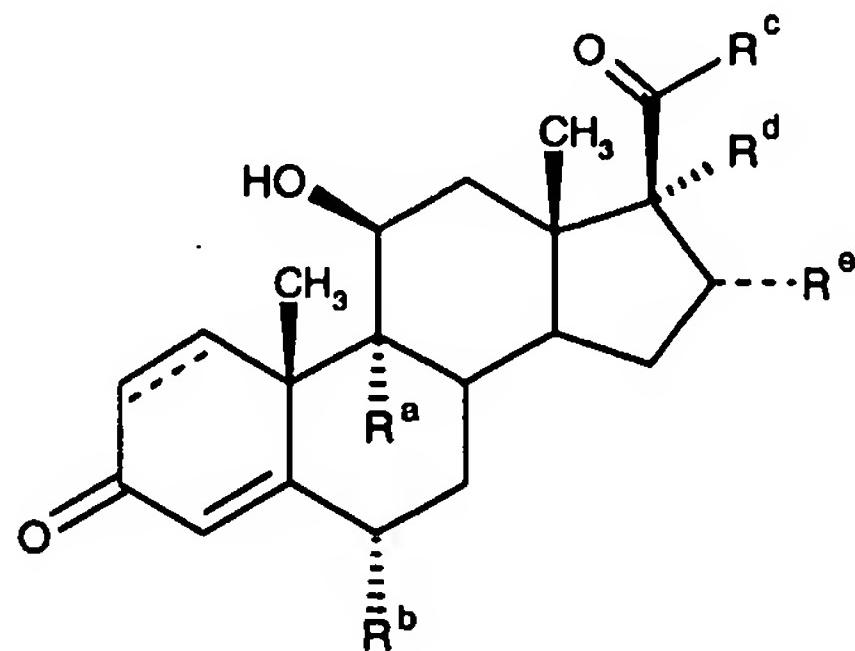
Compounds of Formula I may generally be obtained so that: first the N(CH₃)₂ group of the macrolide subunit is modified, and then the modified macrolide subunit is linked by a chain to the steroid subunit; or, first the steroid and the macrolide subunit possessing the N(CH₃)₂ group are linked by a linking group through another position of the macrolide molecule and then the chemical change on the present N(CH₃)₂ group is performed. The linkage group L may be, for example, attached to the macrolide subunit of Formula VIc at position 9a or 11.

To prevent undesirable side-reactions, it is frequently necessary to protect certain groups such as e.g. hydroxy or amino groups. For this purpose, a large number of protective groups can be employed [Green TW, Wuts PGH, Protective Groups in Organic Synthesis, John Wiley and Sons, 1999] and their selection, use and elimination present the usual methods in chemical synthesis, as is well within the skill for the art..

A still further aspect of the present invention relates to methods for making the compounds represented by Formula I comprising:

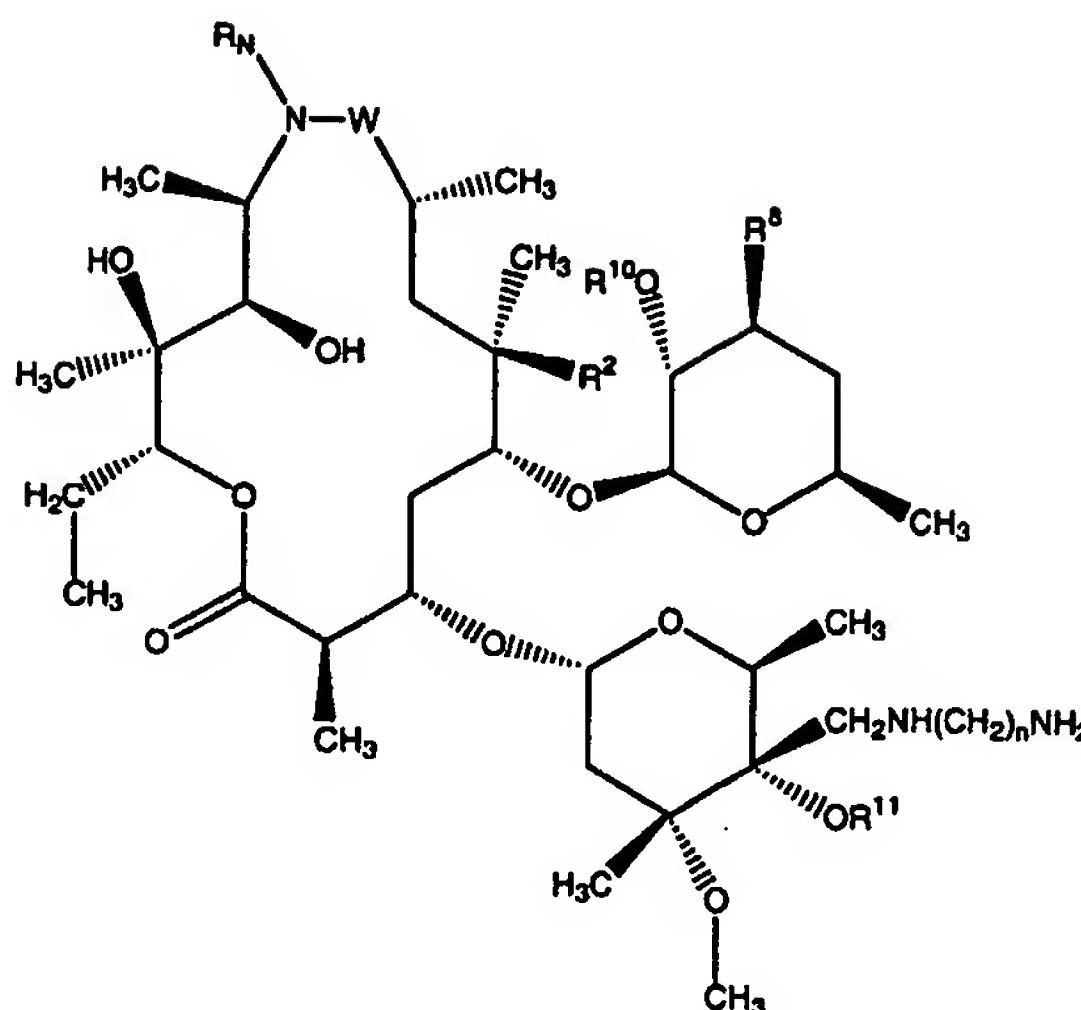
d) for the compounds of Formula I, wherein X¹ is -CH₂- and Q are -NH-(L is VB);

a reaction of (i) the carboxylic acid of the steroid subunit of the structure IVa:



IVa

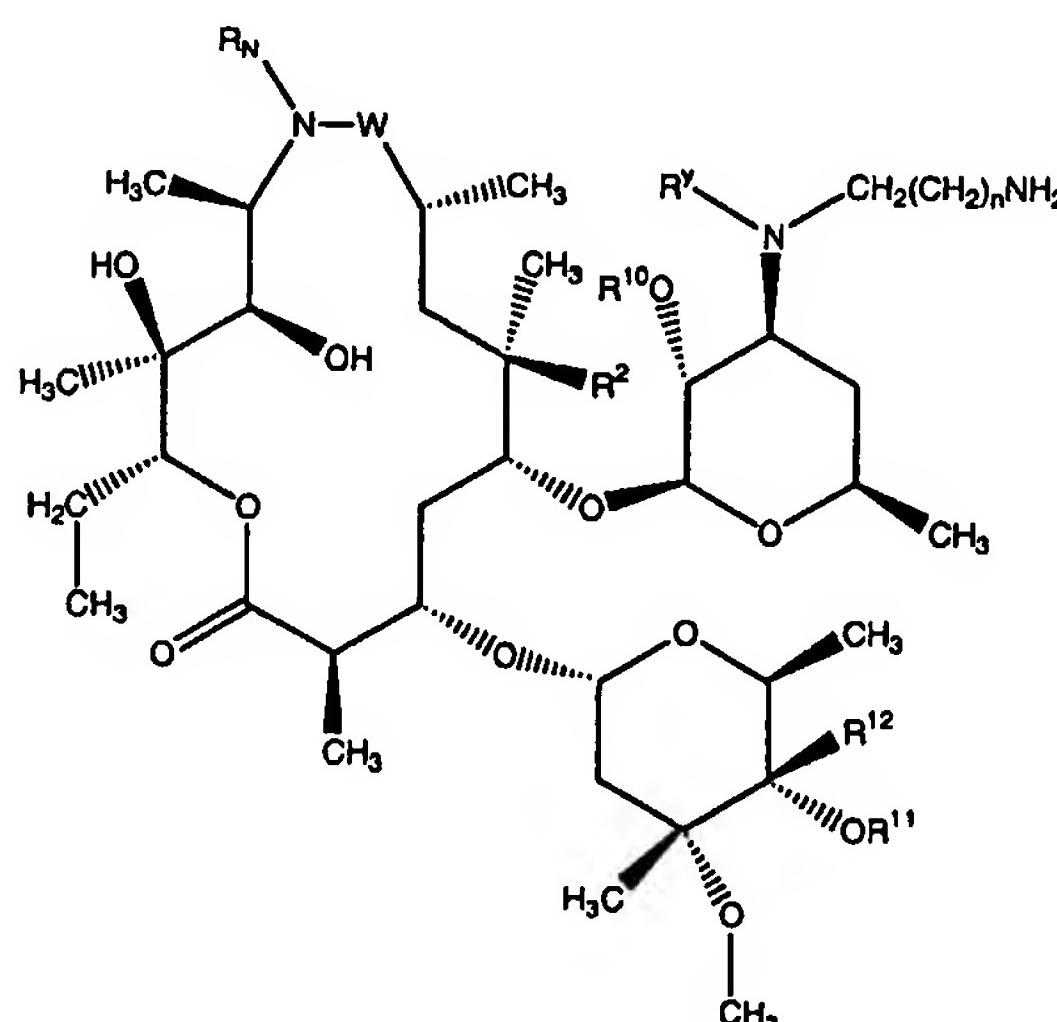
wherein -C(O)R^c is an activated carboxylic acid, and of (ii) a free amino group of the macrolide subunit of the structure Va:

**Va**

wherein W is $>\text{C=O}$ or $-\text{CH}_2-$; R_N, R², R¹⁰, R¹¹, and R⁸ are as defined in Formula II and n is as defined in Formula VA.

e) for the compounds of the structure I, wherein Q and X¹ are $-\text{CH}_2-$ (L is VB),

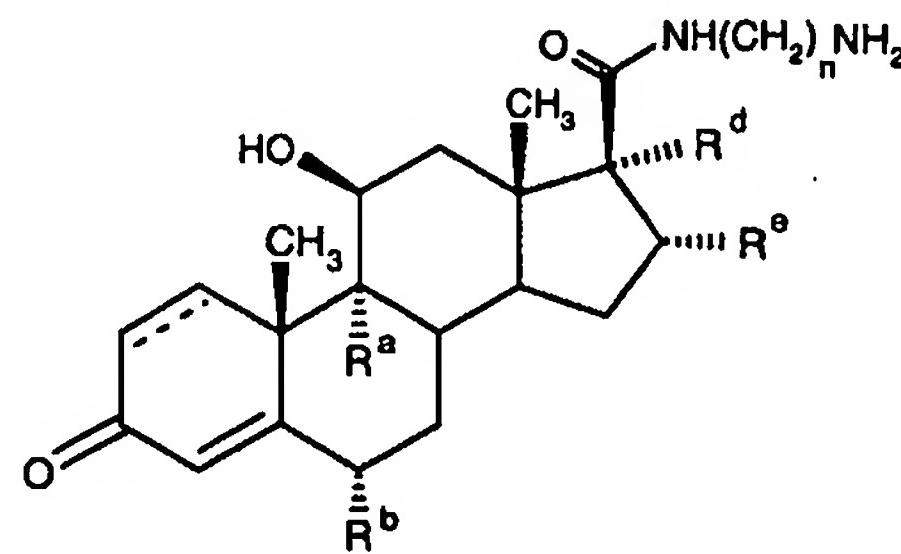
a reaction of carboxylic acid of the steroid subunit of the structure IVa, wherein $-\text{C(O)R}^c$ has the meaning of activated carboxylic acid and of free amino group of the macrolide subunit of the structure Vb:

**Vb**

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wherein W is >C=O or -CH₂-; R_N, R², R¹⁰, R¹¹, R¹², and R^y are as defined in Formula II and n is as defined in Formula VA.

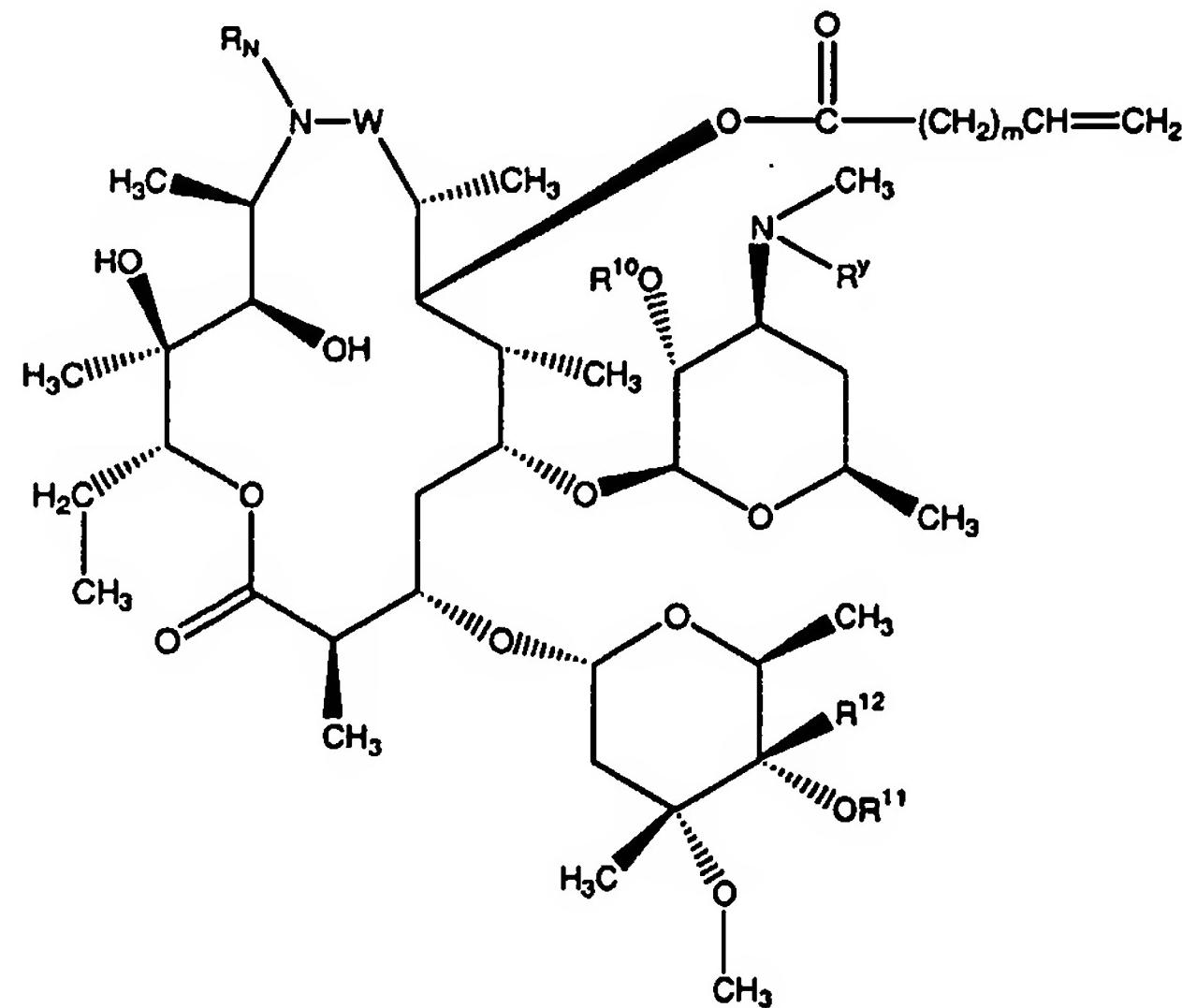
f) for the compounds of Formula I, wherein X¹ is -C(O)- and Q is -NH-, (L is VB),

by reaction of the free amino group of the steroid subunit of the structure IVc:

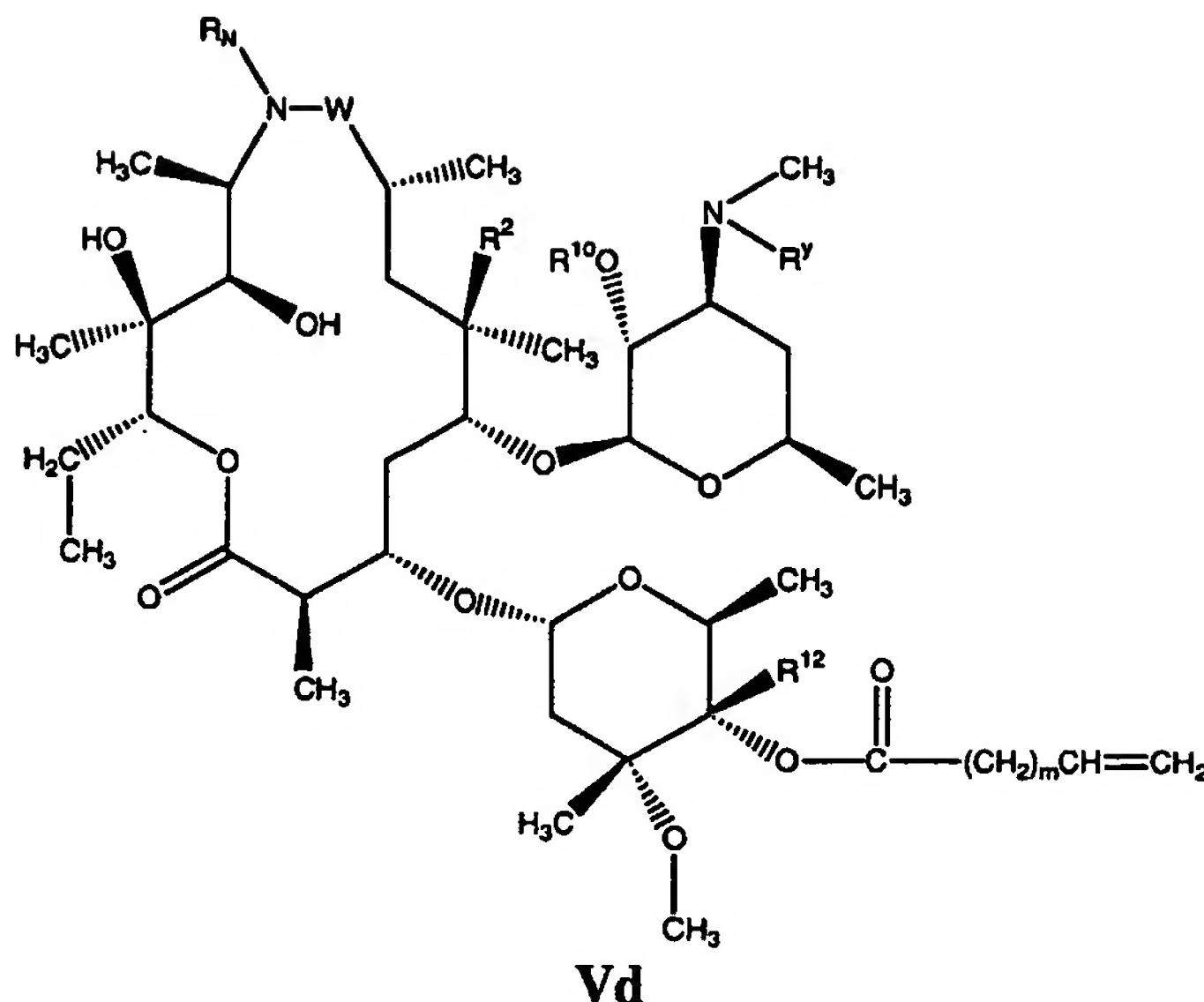


IVc

wherein R^a, R^b, R^d, and R^e are as defined in Formula X, and of a -C=C- bond of the macrolide subunit of the structure Vc or Vd:



Vc

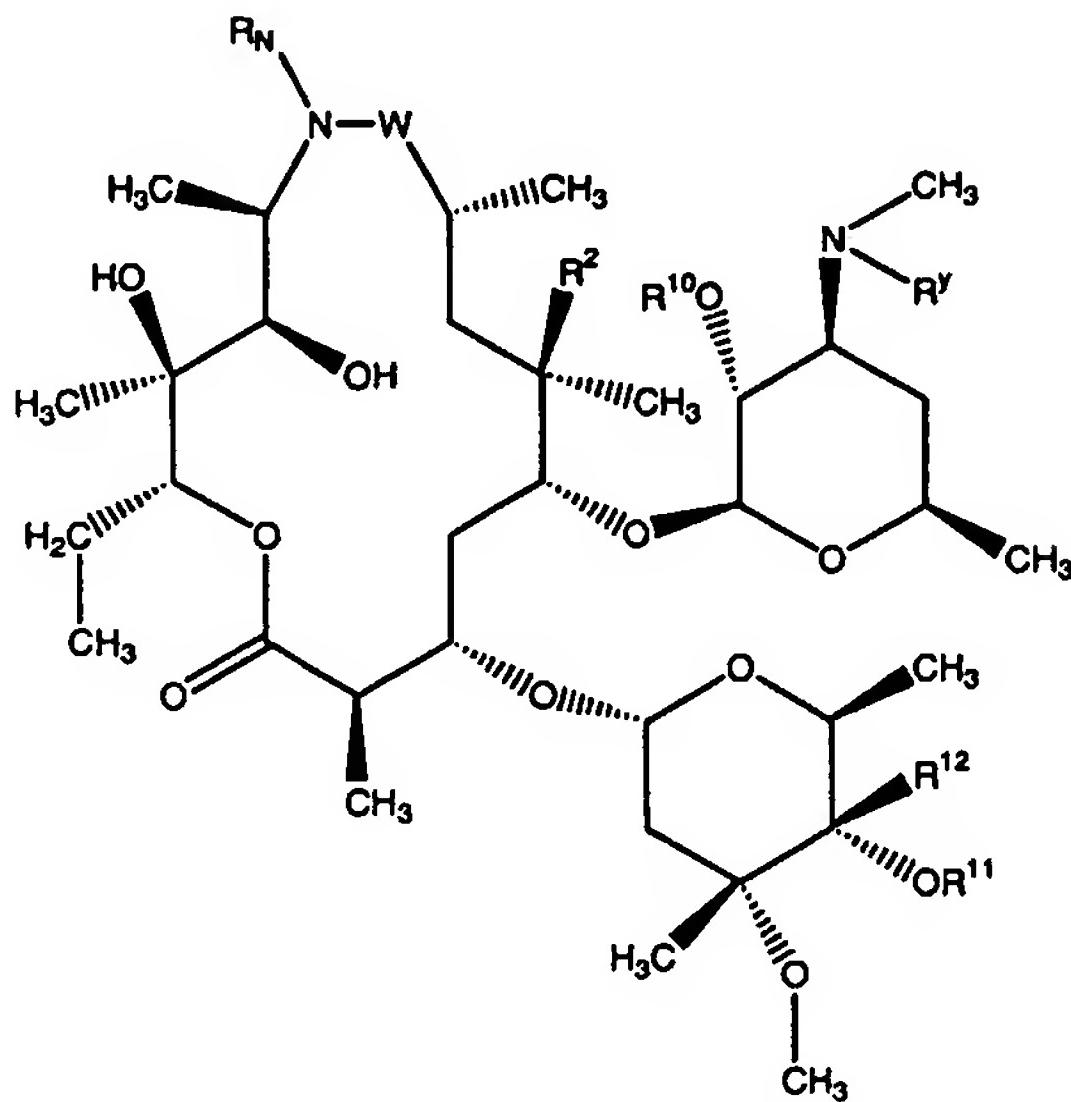


wherein W is >C=O or -CH₂-; R_N, R², R¹⁰, R¹¹, R¹², and R^y are as defined in Formula II and n is as defined in Formula VA.

Methods of Preparation:

d) The compound of Formula I is prepared by a reaction of the carboxylic acid of the steroid subunit of Formula IVa and the amino group of the macrolide subunit of the structure Va, whereby an amide bond is prepared; it is carried out using the usual acid derivatives having an activating effect on carboxylic acid, such as halogenides, mixed anhydrides especially carbodiimides or benzotriazole. The reaction proceeds in the presence of a base (mainly organic) e.g. triethylamine at room temperature under an inert atmosphere, such as nitrogen or argon over a period of several hours to several days (Romo D et al., *J. Am. Chem. Soc.* **1998**, *120*:12237). The steroid subunits of the structure IVa are either commercially available products or have been obtained by the described methods (Suzuki T et al., *Chem. Soc., Perkin Trans. I* **1998**, 3831-3836; McLean HM et al., *J. Pharm. Sci.* **1994**, *83*:476-480; Little RJ et al., *Pharm. Res.* **1999**, *16*:961-967; Kertesz DJ et al., *J. Org. Chem.* **1986**, *51*:2315-2328 and Patent Application US 471049).

The starting macrolide subunits of the structure **Va** may be obtained by the action of the corresponding alkylene-diamines on the macrolide subunits of the structure **VI**:

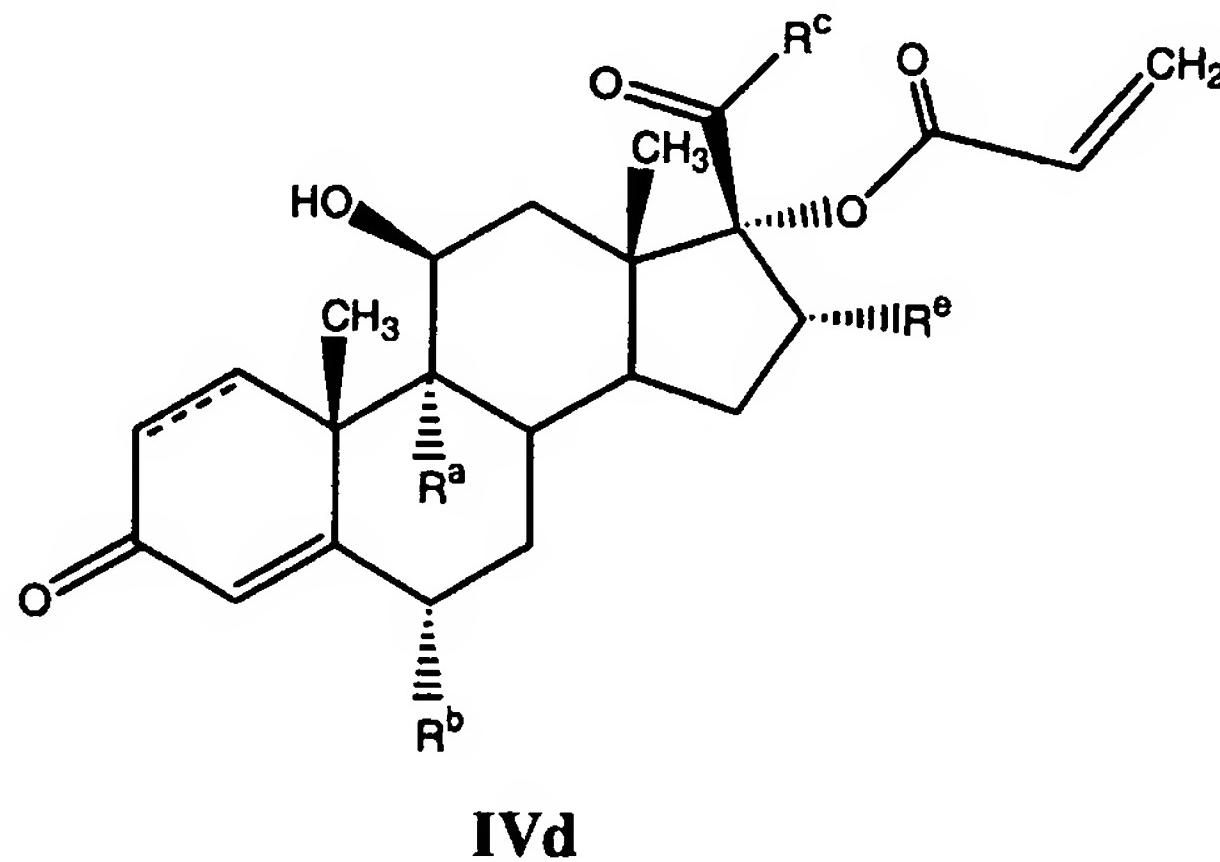
**VI**

wherein R^{11} and R^{12} together with the adjacent C atom from the cladinosyl ring form a three-membered oxirane ring. Such compounds were prepared by the procedures described for the preparation of analogue compounds in Patent Application WO 98/56801. Thus, for example, by oxidation of the free hydroxyl group OR^{11} , where R^{11} is H, of the macrolide subunit of the structure **VI**, a corresponding ketone is obtained, where R^{11} and R^{12} form a $>C=O$ group with the C/4" carbon atom which, by reduction with a metal hydride (preferably NaH) under trimethylsulphoxonium-iodide yields the above mentioned oxirane ring.

e) The preparation of a compound within Formula **I** is carried out by a reaction of the carboxylic acid of the steroid subunit of Formula **IVa** and a free amino group of the macrolide subunit of Formula **Vb**, whereby an amide bond is prepared, and the process proceeds in the manner and under conditions as described in the method d) above. The starting macrolide subunits of the structure **Vb** may be obtained by the action of a corresponding cyanoalkylhalogenide or cyanoalkenyl (preferably acronitrile) on the macrolide subunits of the structure **VI**, wherein R^y is H, and which may be obtained by demethylation of the macrolide subunit of the structure **VI** wherein R^y is a CH_3 group under UV radiation (Bright M et al., *J. Antibiot.* 1988, 41:1029).

f) A compound within Formula I may be prepared by reacting of an amino group of the steroid subunit of Formula IVb and the -C=C- group of the macrolide subunit of the structure Vc or Vd, whereby an amino bond is prepared. Steroid subunits of the structure IVb are either known compounds or may be obtained from the steroid subunit IVa, wherein R^c is an OH group, by the action of alkylene-diamine. The starting macrolide subunits of the structure Vc and Vd may be obtained by the action of a corresponding halogenalkanoylchloride (preferably 3-chlorpropionylchloride) on the free OH group of the macrolide subunit of Formula VI.

g) for the compounds of Formula I, wherein X¹ is -C(O)- and Q is -NH-, (L is VB), by reaction of a -C=C- bond of the steroid subunit of the structure IVd:



wherein R^a, R^b, R^c, and R^e are as defined in Formula X, and the free amino group of a macrolide subunit of the structure Va, VIa, VIb or Vb that are described before.

Methods of Preparation:

g) The compound of Formula I is prepared by a reaction of the steroid subunit of Formula IVd and the amino group of the macrolide subunit of the structure Va, VIa, VIb or Vb whereby an amide bond is prepared. The starting steroid subunits of the structure IVd may be obtained by the action of a corresponding halogenalkanoylchloride (preferably 3-chlorpropionylchloride) on the steroid subunit

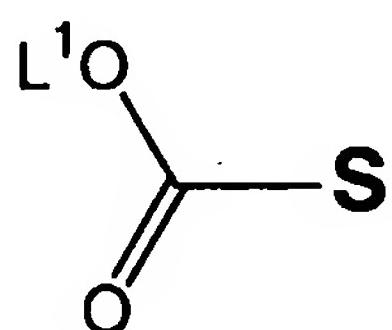
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of Formula IVa wherein R^d is an OH group (Phillipps, G. et al., J. Med. Chem. 1994, 37, 3717-3729).

Compounds of Formula I may generally be obtained in the following way: one end of the linker chain L is first linked to the macrolide subunit M, and then the other end of the chain is joined to the steroid subunit; or, one end of the chain L is first linked to the steroid subunit S and then the other end of the chain to the macrolide subunit M, or finally, one part of the chain is linked to the macrolide subunit M, whereas the other part of the chain is linked to the steroid subunit S, with the ends of the chain parts being then chemically linked to form the chain L.

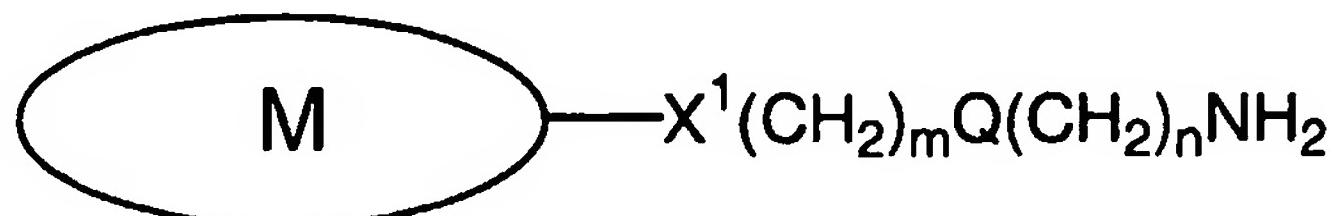
More general schemes for making the compounds of the invention are apparent to a person of skill in the field of the invention in light of the foregoing. Compounds within Formula I can be prepared by the following processes.

- a) Compounds of Formula I, where X² is -NH-, can be formed by reacting (i) a steroid anti-inflammatory subunit represented by Formula V:



V

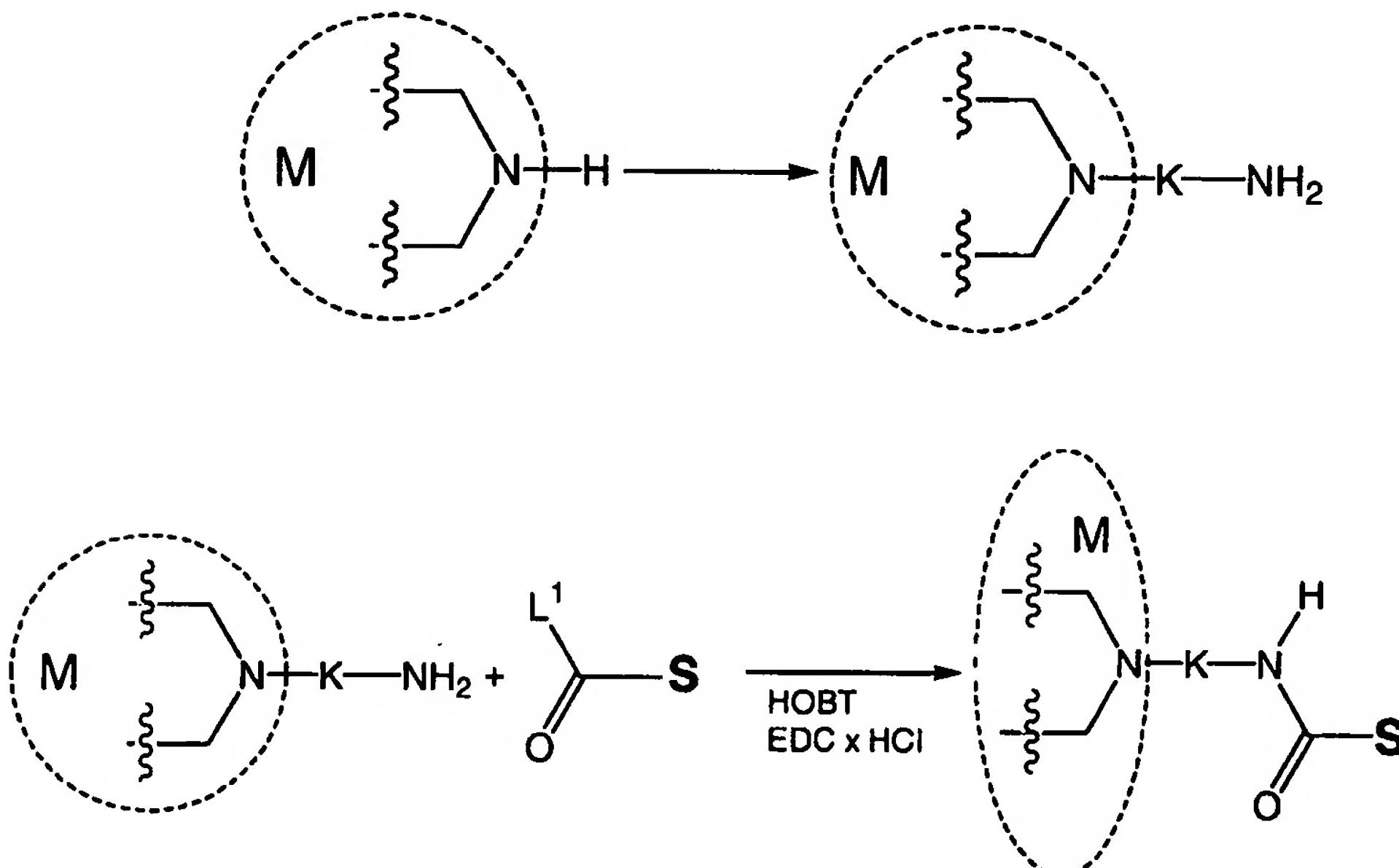
wherein L₁ represents a leaving group (such as hydroxy), and (ii) a free amino group of a macrolide subunit represented by Formula VIId:



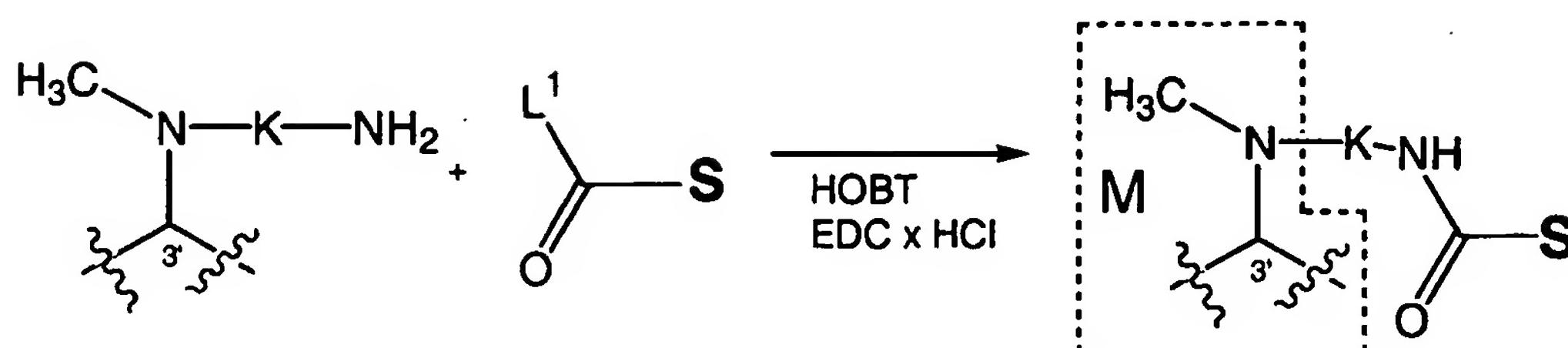
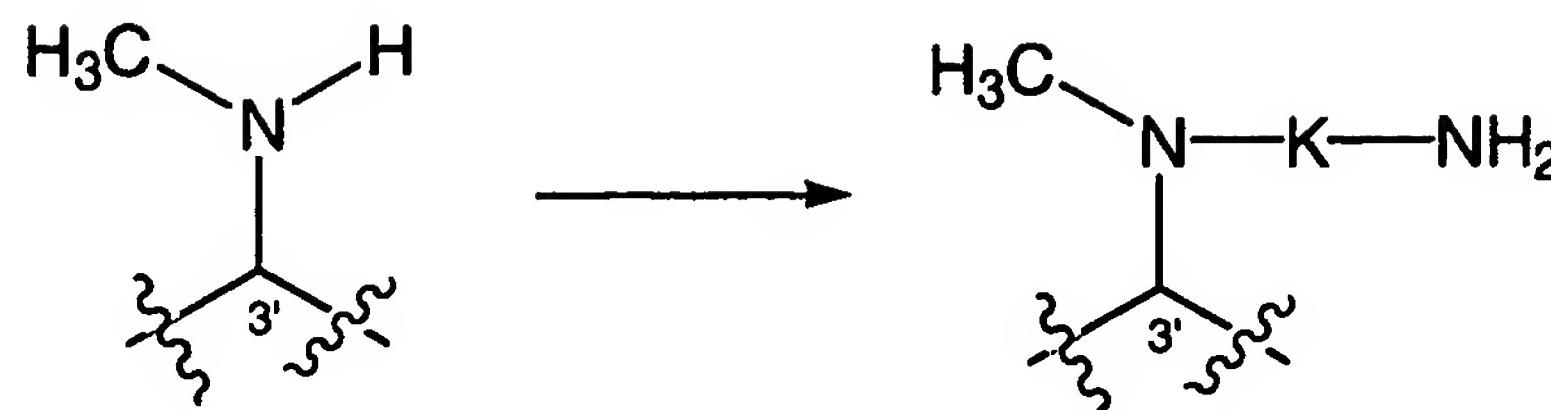
VIId

The reaction is generally performed with acid derivatives which have the ability to activate the carboxylic acid group of steroidal anti-inflammatory subunit, such as halogenides, mixed anhydrides and especially carbodiimides (such as -(3-dimethylaminopropyl)-3-ethyl-carbodiimide (EDC)) and benzotriazoles. The reaction proceeds in the presence of a base, such as an organic base (e.g., triethylamine), at room temperature under an inert atmosphere such as nitrogen or argon. The reaction may require several hours to several days to come to completion.

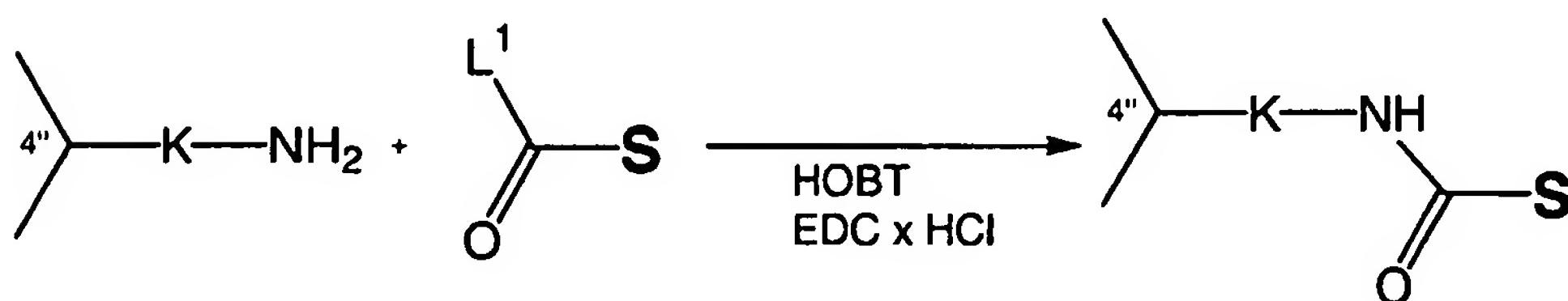
For example, when L is -K-NH- (wherein K is the portion of the L molecule attached to the macrolide) the compound of Formula I can be formed by derivatizing an NH group on the macrolide ring to an -N-K-(NH₂)- group and reacting the derivatized macrolide with a steroid anti-inflammatory subunit represented by Formula V:



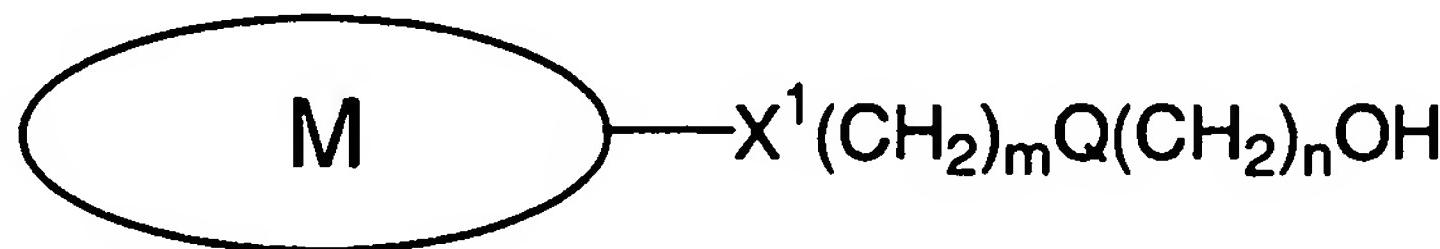
This process may also be performed when the NH group in the macrolide is attached at the 3' position of a sugar ring S¹ (i.e., a desozamine sugar) of the macrolide:



or the 4 position of the sugar ring S²:



b) Compounds represented by Formula I, where X² is -OC(O)-, can be formed by reacting a compound of Formula V and the free hydroxyl group of a macrolide subunit represented by Formula VIe:



VIe

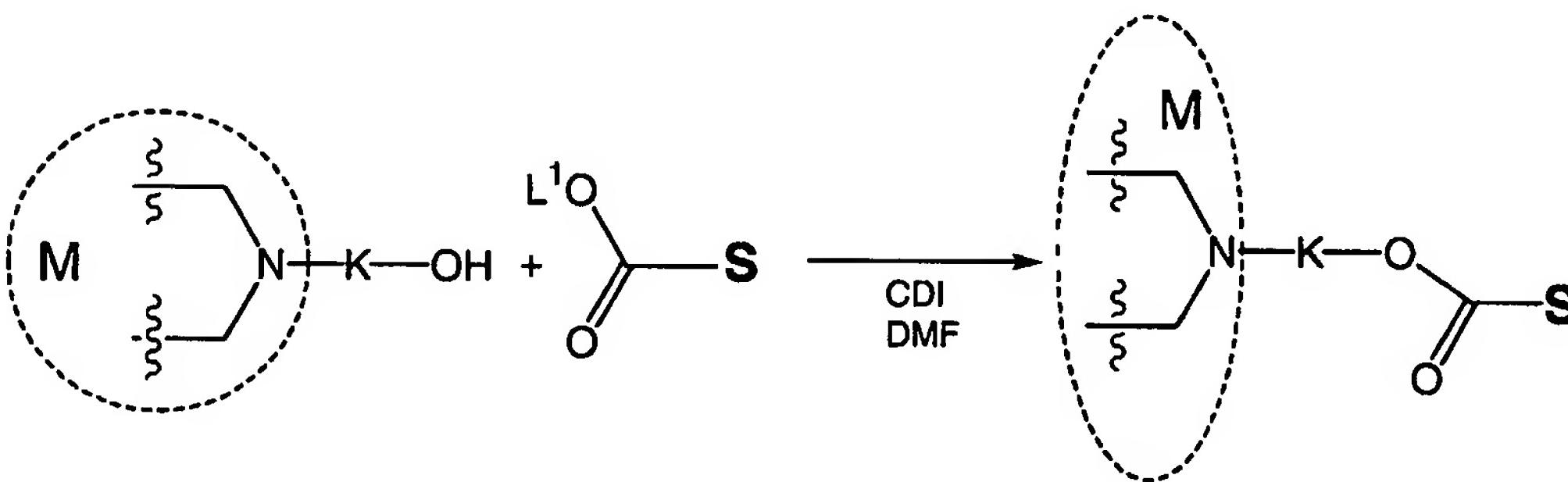
The reaction is generally performed with acid derivatives which have the ability to activate the carboxylic acid group such as halogenides, mixed anhydrides and especially carbodiimides and benzotriazoles. The reaction is typically performed

at room temperature under an inert atmosphere such as nitrogen or argon.

The reaction may require several hours to several days to come to completion.

The starting macrolide subunits of the structure **VIIb** are known compounds or may be obtained according to the procedures described for analogous compounds, such as those described in Costa AM et al., *Tetrahedron Letters* 2000, 41:3371-3375, which is hereby incorporated by reference in its entirety. See, also Bright U.S. Patent No. 4,474,768 and Bright, G.M., et al, *J. Antibiot.*, 1988, 41:1029-1047, both incorporated in their entirety by reference.

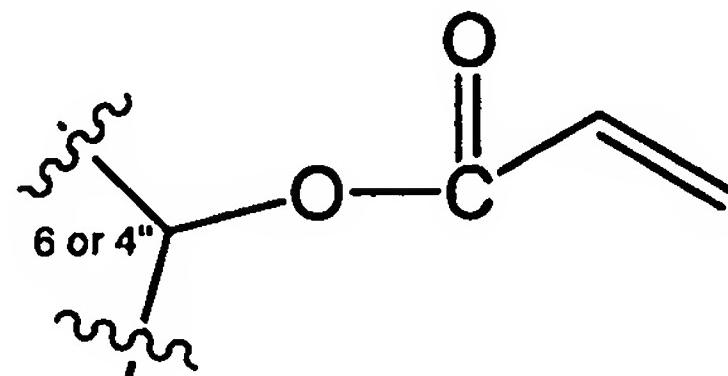
For example, when linkage L is -K-O-, the compound of Formula I can be formed by (1) derivatizing an NH group on a macrolide to an N-K-OH group and (2) reacting the derivatized macrolide with the free carboxylic acid group on a steroid S:



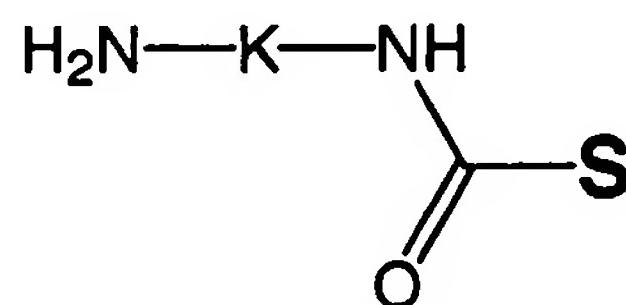
The linkage group -K-OH can be attached to the secondary nitrogen atom of the macrolide subunit as follows. The macrolide subunit is reacted with an alkenoyl derivative, such as $\text{CH}_2=\text{CH}(\text{CH}_2)_m$, $\text{C}(\text{O})\text{O}\text{-Alkyl}$ (e.g., methylacrylate). The ester group (i.e., $-\text{C}(\text{O})\text{O}\text{-Alkyl}$) is then reduced, such as with a metal hydride (e.g., LiAlH_4) in an anhydrous organic solvent, to yield the macrolide subunit having the linkage group -K-OH (i.e., M-K-OH). The reduction is typically performed at a low temperature and preferably at 0° C or lower.

This process can also be performed when the NH group is attached at the 3' position of a sugar ring in the macrolide (such as a sugar at the 3 position of the macrolide).

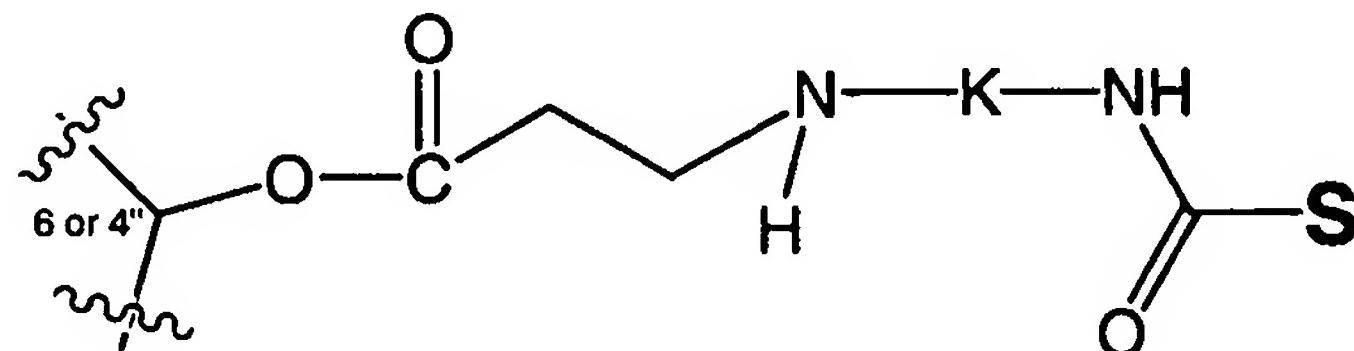
c) Compounds represented by Formula I, wherein X^1 is $-OC(O)-$, Q is $-NH-$ and X^2 is $-NH-$ can be prepared by reacting a macrolide subunit represented by the formula



For example this process can be performed when OH group is attached at the C/6 or C/4" position of the macrolide subunit represented by Formula:.

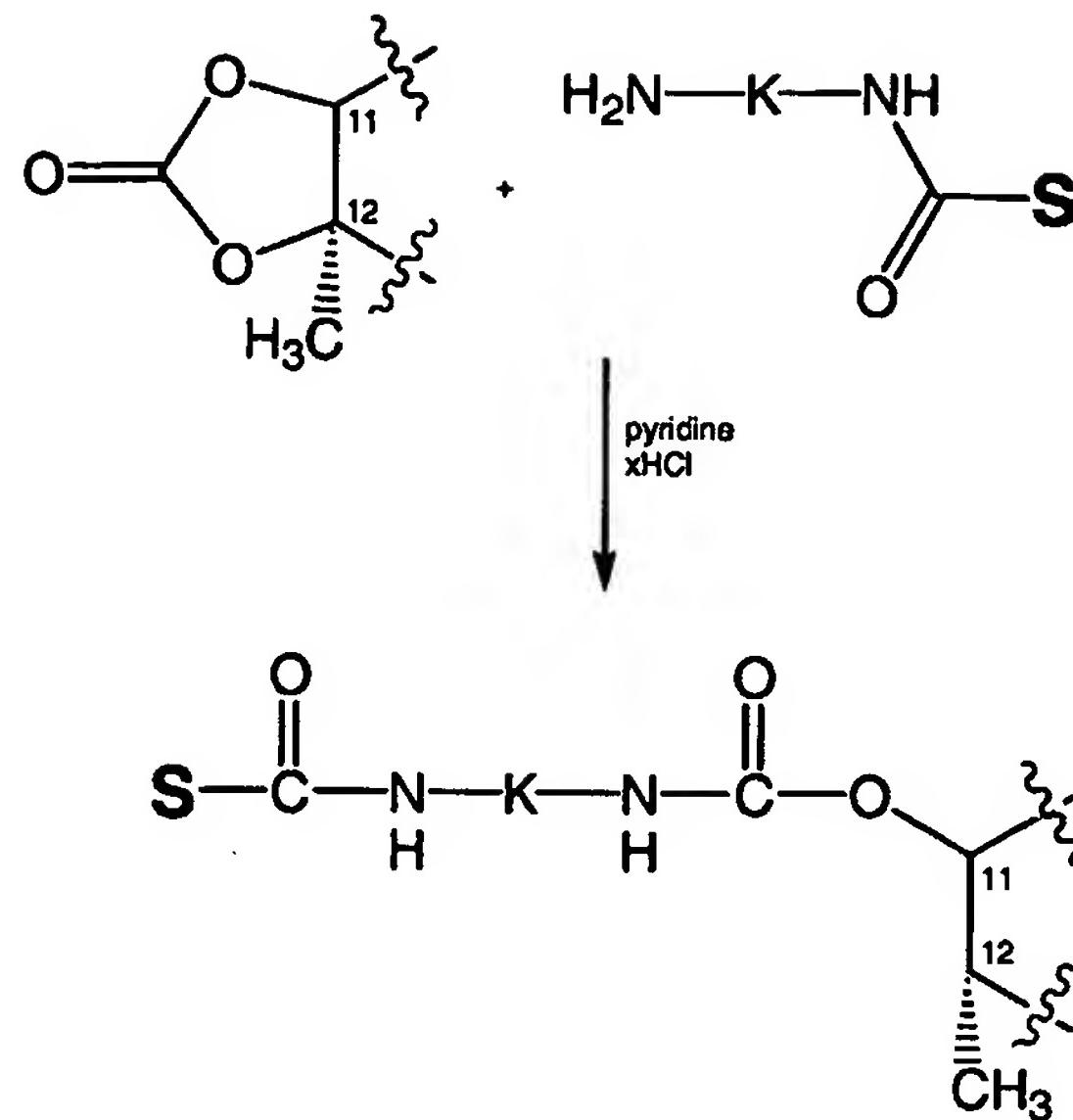


in a solvent, such as acetonitrile, to yield

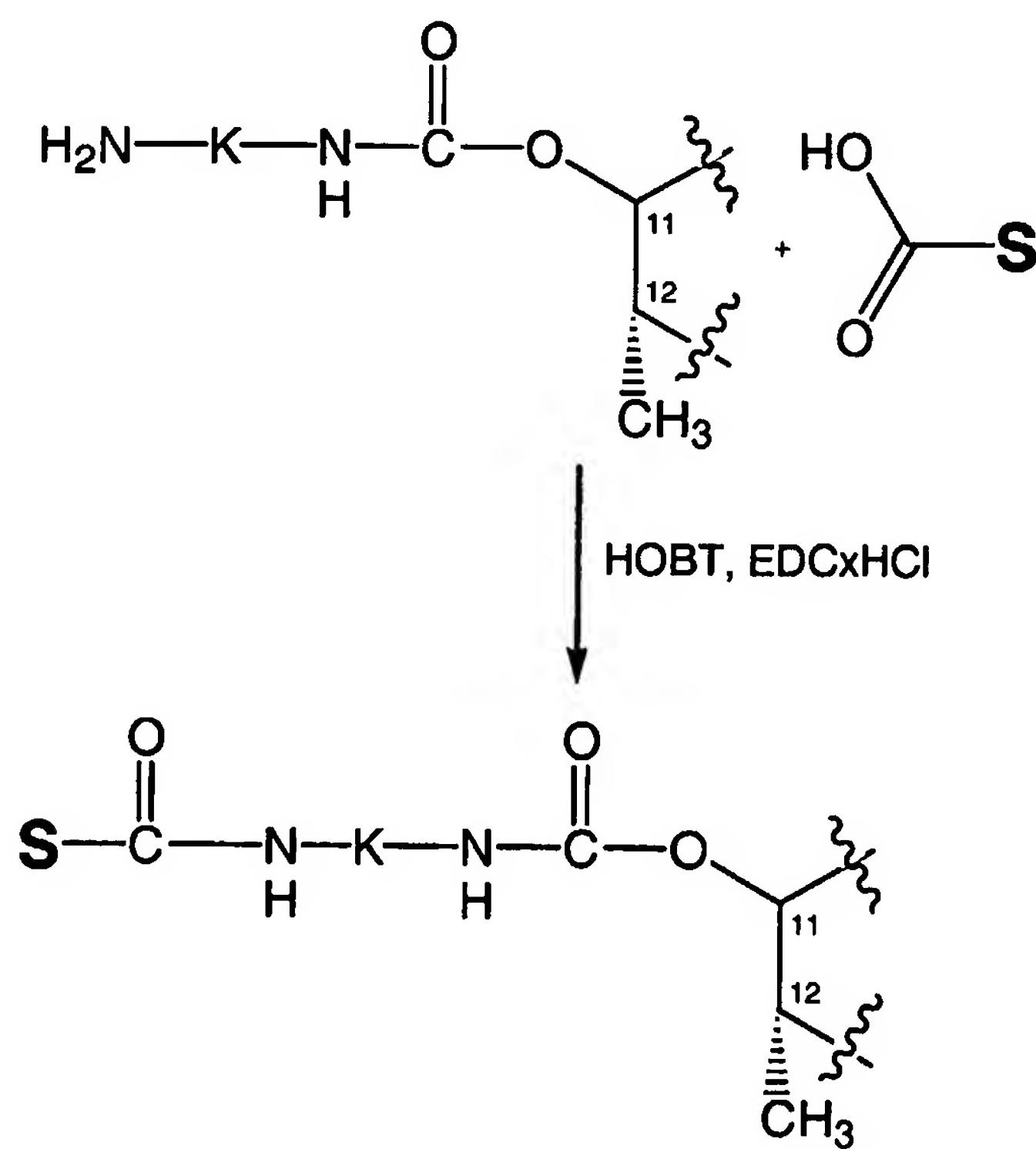


The derivatized steroid (i.e., $S-C(O)-NH-K-NH_2$) may be formed by reacting an appropriate amine (having the linkage group $-K-NH_2$) with a carboxylic acid group or an ester group of a steroid according to Formula V.

d) Compounds represented by Formula I, where X^2 is $-NH-$, can be prepared by reacting a macrolide subunit and a derivatized steroid subunit having a free amino group as shown below.



e) Compounds represented by Formula I, where X^2 is -NH-, can be prepared by reacting a macrolide subunit and a steroid subunit having a free carboxylic acid group as shown below.



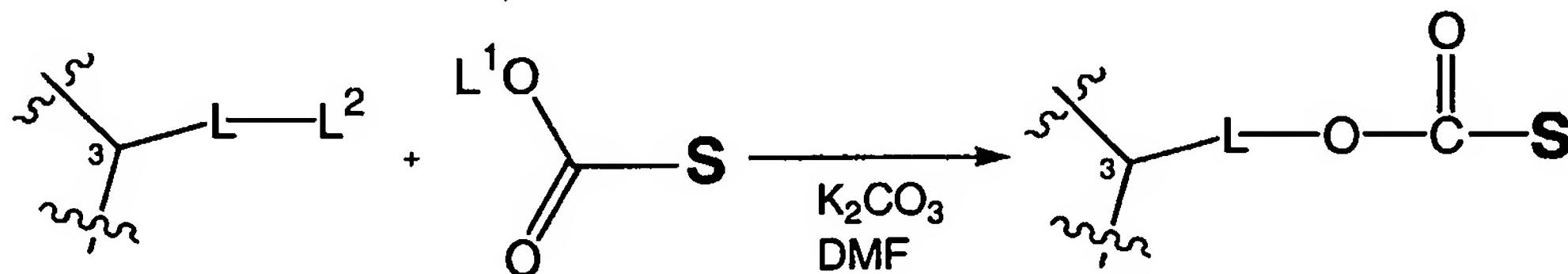
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The reactant macrolide subunit can be formed by oxidizing the corresponding macrolide having a hydroxy substituent at the "4" position on

cladinose sugar to obtain a =O substituent at the "4" position, converting the

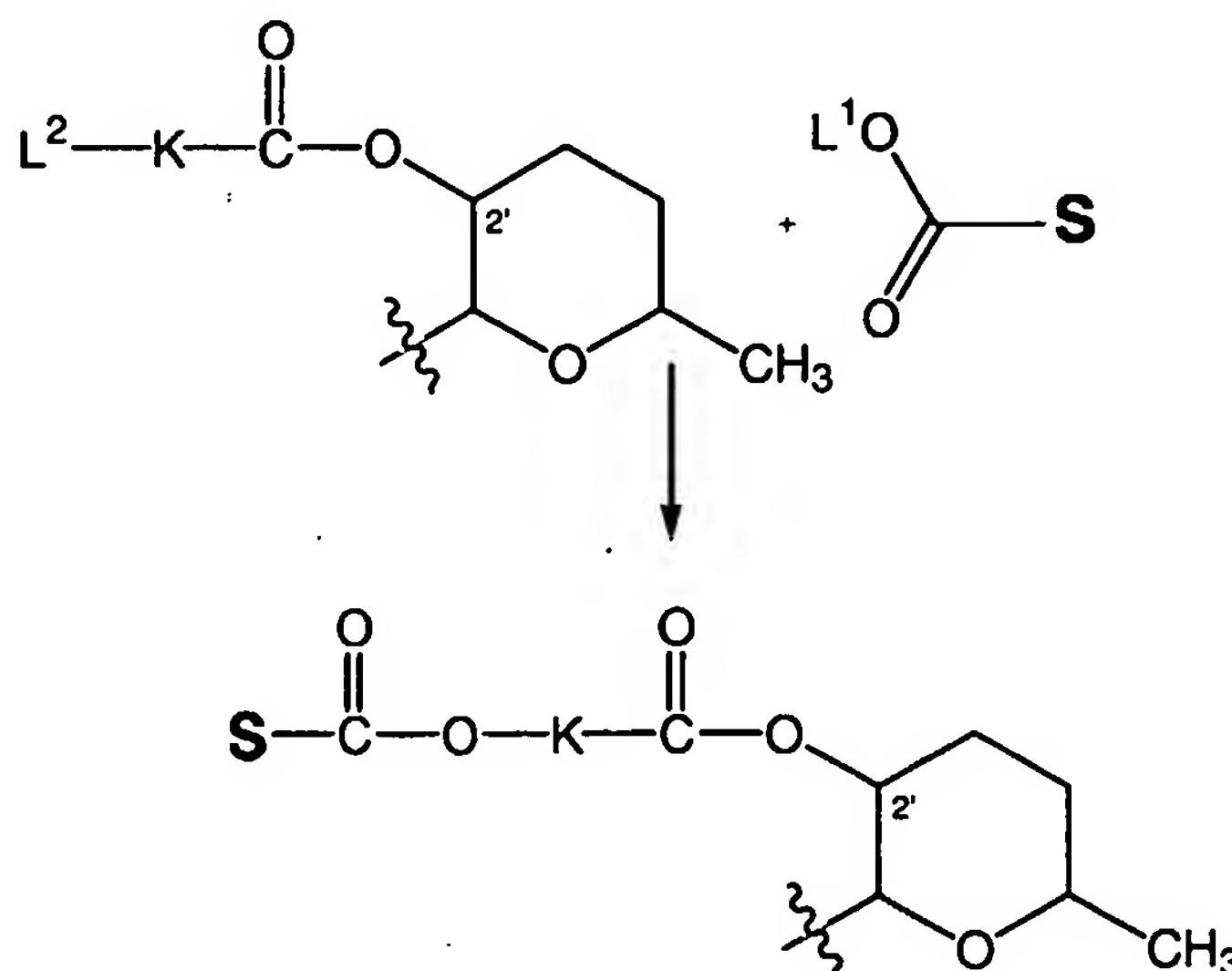
at the "4" position to an epoxy group (), and cleaving the epoxy group with an appropriate reactant(s) to yield the macrolide subunit (M-O-C(O)-NH-K-NH₂).

f) Compounds of Formula I can be prepared by reacting a macrolide subunit having a leaving group L² (such as Br), and a steroid as shown below.



The starting macrolide subunit can be prepared by cleaving the sugar group attached at the 3-position of the macrolide ring and then reacting the macrolide with a reagent of Formula L²-L-L¹, where L² is a leaving group.

g) Compounds of Formula I can be prepared by reacting a macrolide subunit having a leaving group L² (such as Br), and a steroid as shown below.

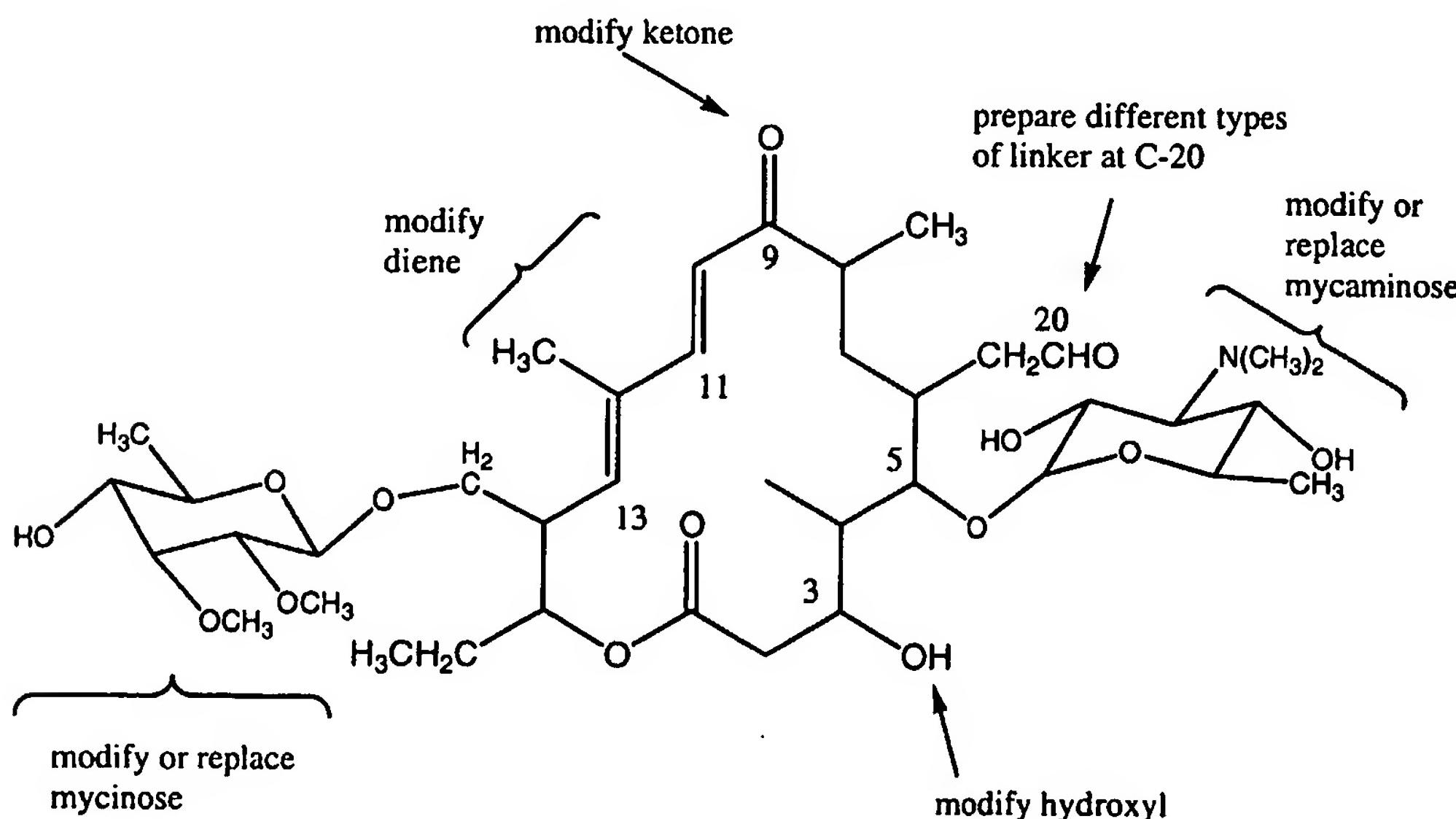


The salts of the compounds represented by Formula I may be prepared by applying generally known procedures such as, e.g., a reaction of the compounds of the structure I with a corresponding base or acid in a suitable solvent or mixture of solvents e.g. ethers (diethyl ether) or alcohols (ethanol, propanol or *iso*-propanol).

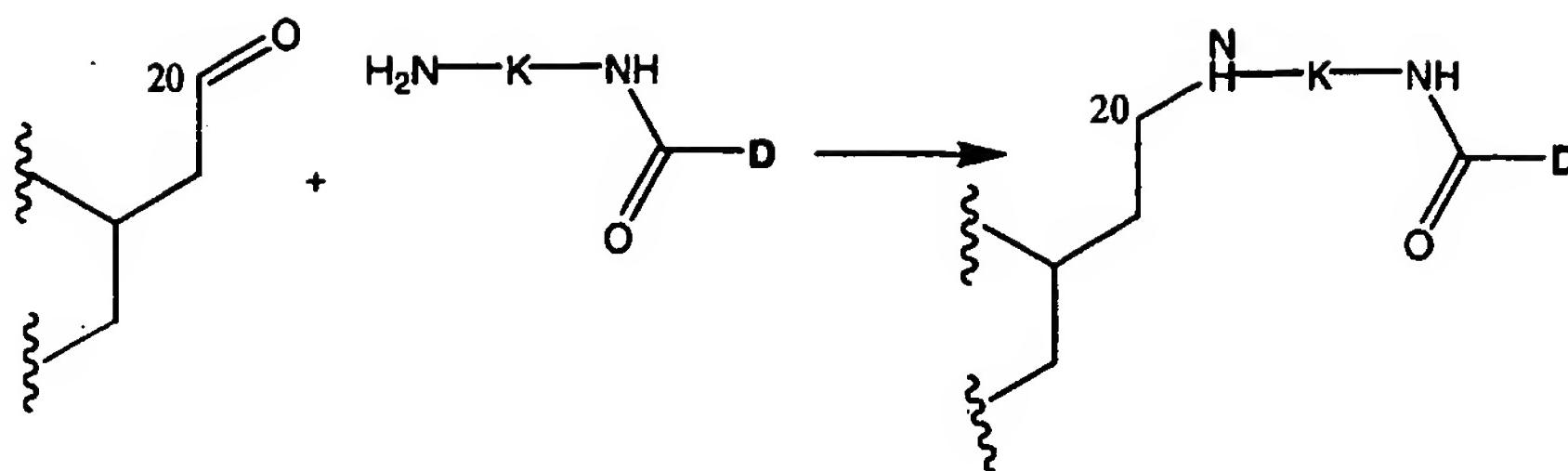
The 16-membered ring macrolides are traditionally divided into sub-families based upon the substitution patterns of their aglycones. The principal prototypes of this family can be represented by leucomycin, spiramycin and tylosin.

Tylosin is a representative of 16-membered macrolides, which possesses a highly substituted aglycone with two double bonds (tylonolide) and a third saccharide substituent (β -D-mycinose) in addition to the disaccharide attached to the 5-hydroxyl group. Hydrolysis of mycarose from disaccharide yielded desmycosyl-tylosin (desmycosin).

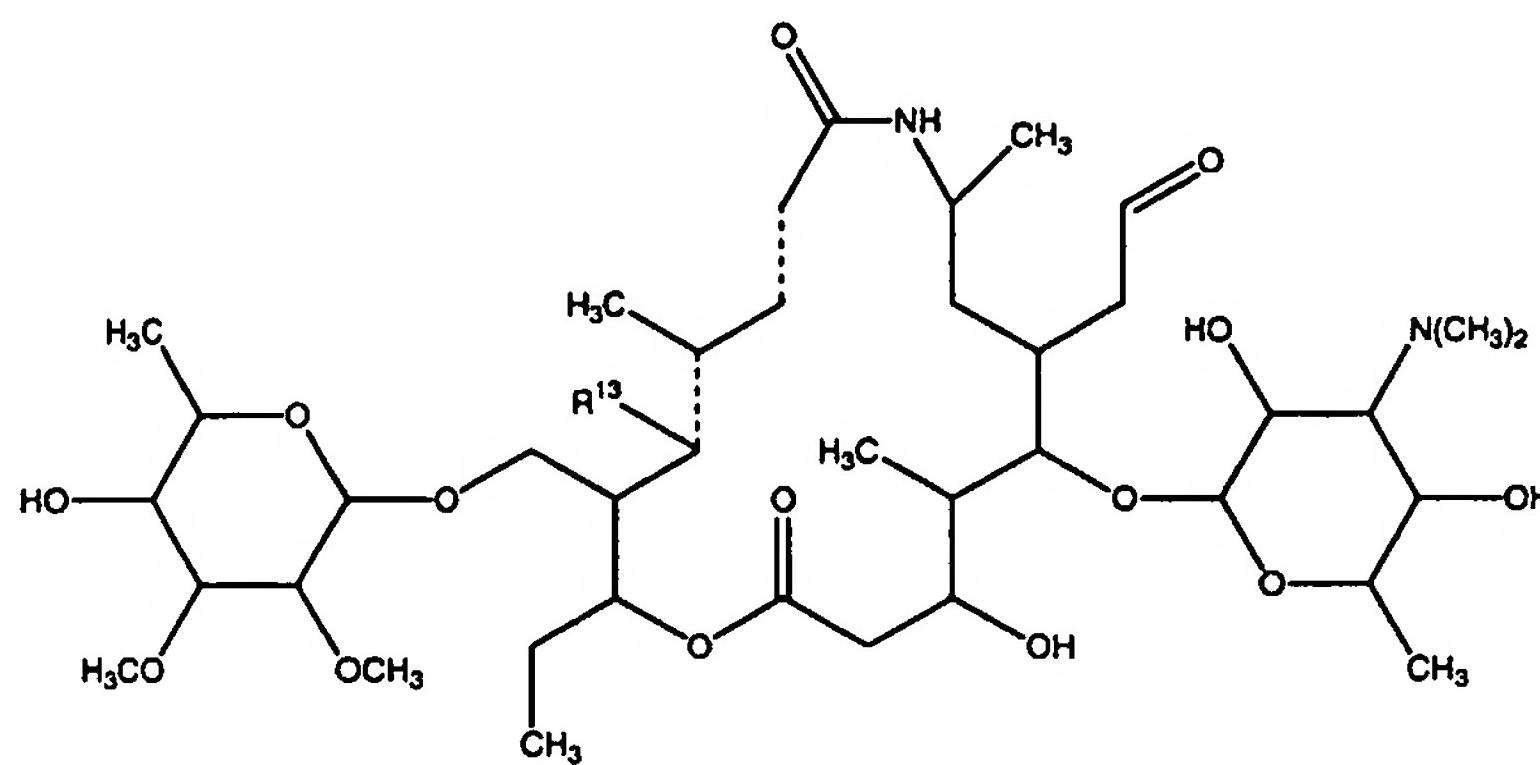
Potential sites of modification in desmycosin:



For example, 16-membered ring macrolide hybrid could be prepared by reductive amination of the C-20 aldehyde group.



This reaction could be used also for 17-membered azalides like 8a-aza-homodesmycosins and its derivatives (such as di- and tetrahydro derivatives).



----- represents a single or double bond

R¹³ is hydrogen or hydroxy

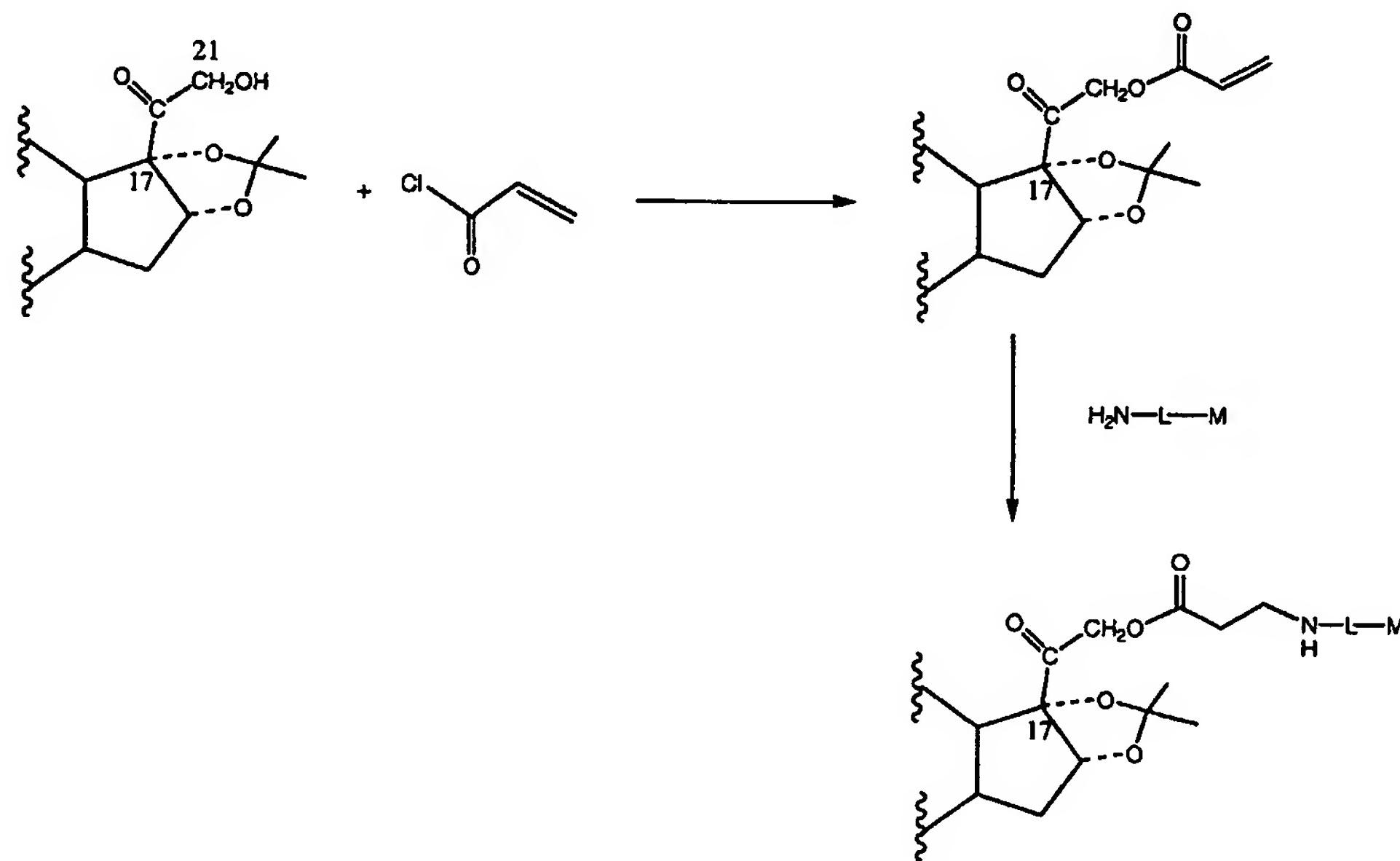
Alternatively, 16-membered ring macrolide derivatisation can proceed by transforming double bonds (e.g., by epoxidation), and cleaving the epoxy group with an appropriate reactant (such as a diamine) to yield the reactant macrolide subunit (M-O-C(O)-NH-K-NH₂).

Also a ketone in position 9 may be modified by hydroxylamine hydrochloride to yield oxime and then reduced to amine.

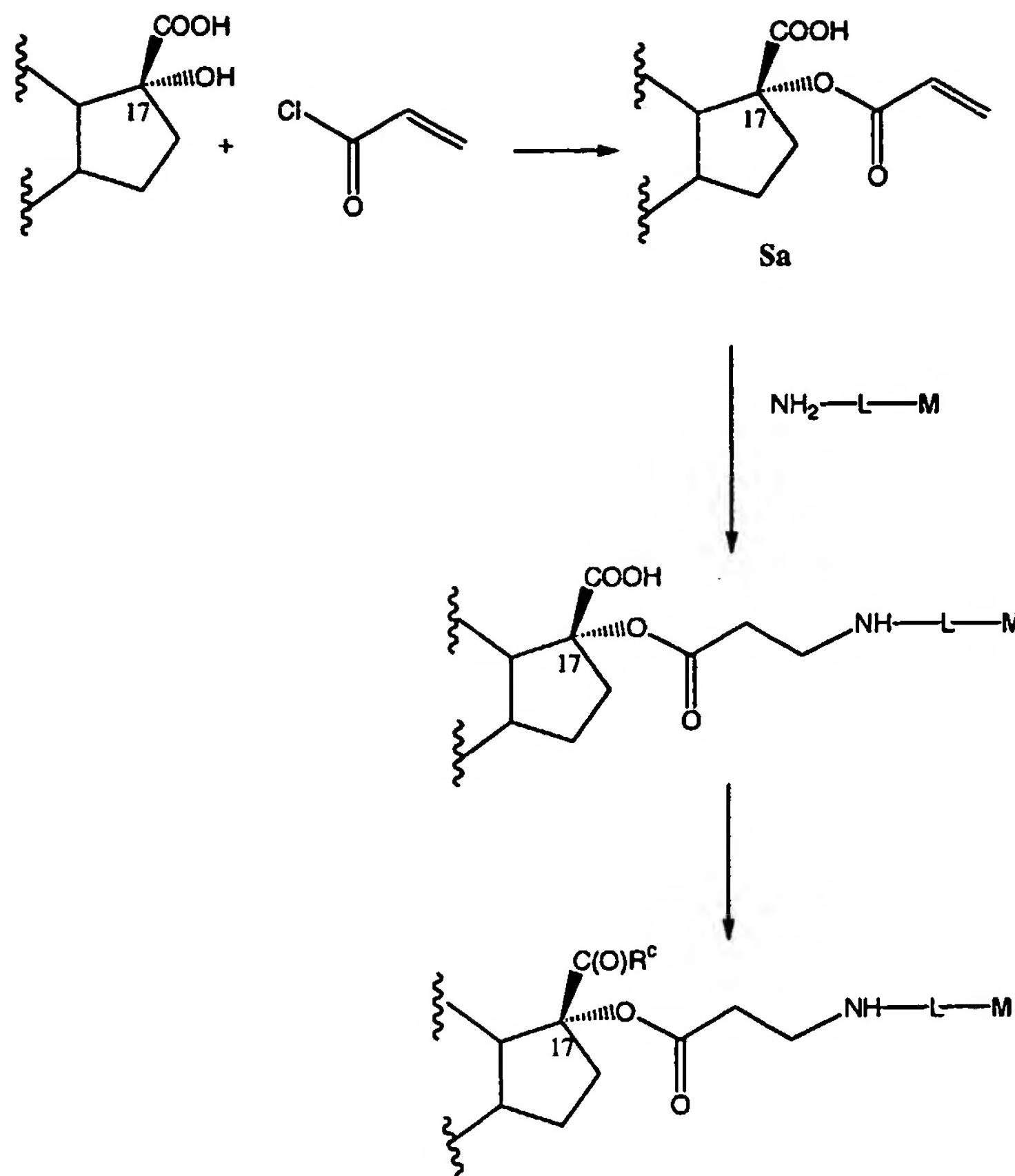
If the steroid is an -S(O)_n-R_C at position 17, the same linkage schemes can be used.

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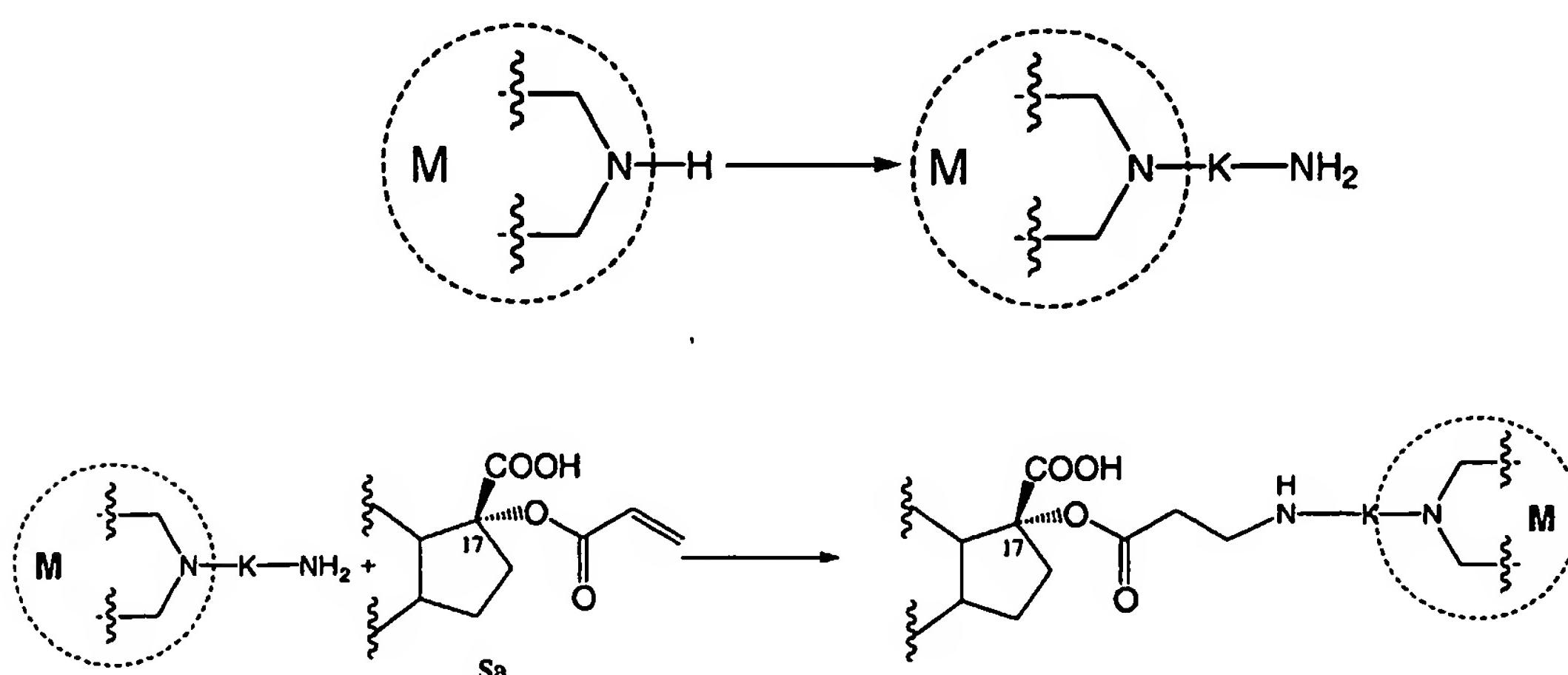
The steroid subunit may be linked to the macrolide through the 21 hydroxy group in steroids that have such a group. Beginning with a 21-hydroxy steroid cyclic ketal is reacted with an appropriate carboxylic acid halide or an anhydride, preferably in a solvent such as methylene chloride in the presence of a tertiary amine base or pyridine at a reduced temperature (-5⁰C to 30⁰C). The intermediate so produced is reacted with H₂N-L-M to form compounds of Formula I



The steroid subunit may also be linked to the macrolide through the 17 position on the steroid subunit. One method for preparing such a compound is as follows:

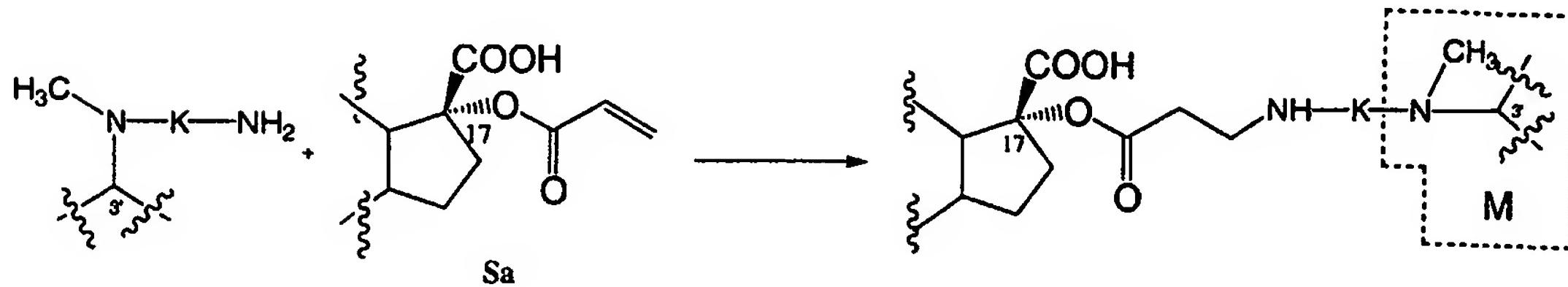
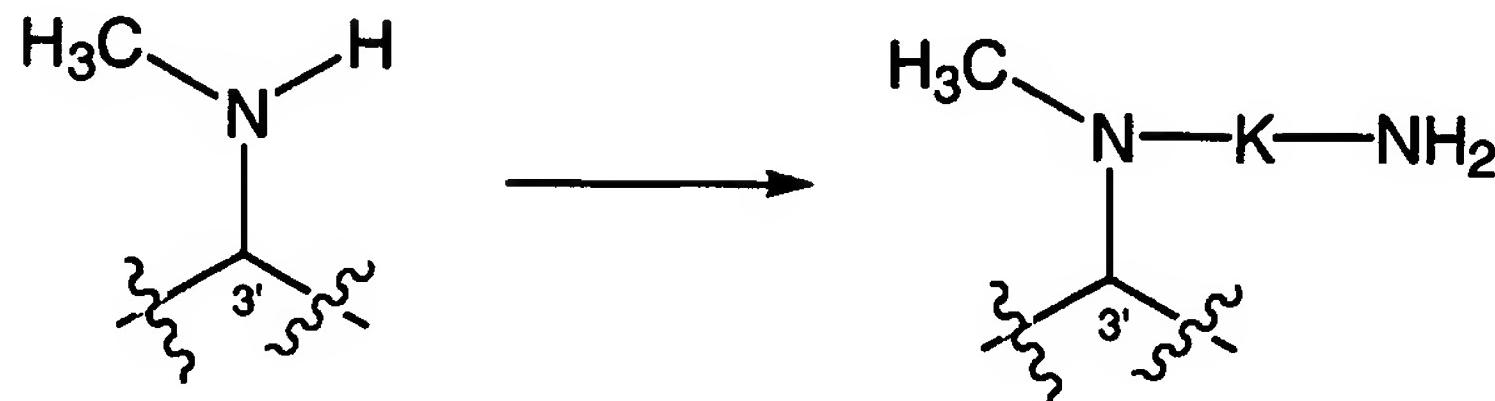


For example, when L is $-\text{K-NH-}$ (wherein K is the portion of the L molecule attached to the macrolide) the compound of Formula I can be formed by derivatizing an NH group on the macrolide ring to an $-\text{N-K-(NH}_2\text{)-}$ group and reacting the derivatized macrolide with a steroid anti-inflammatory subunit represented by Formula **Sa**:

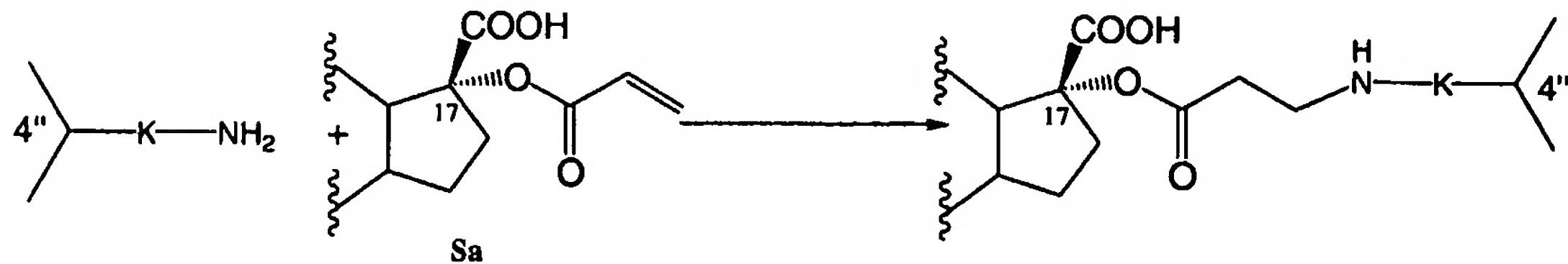


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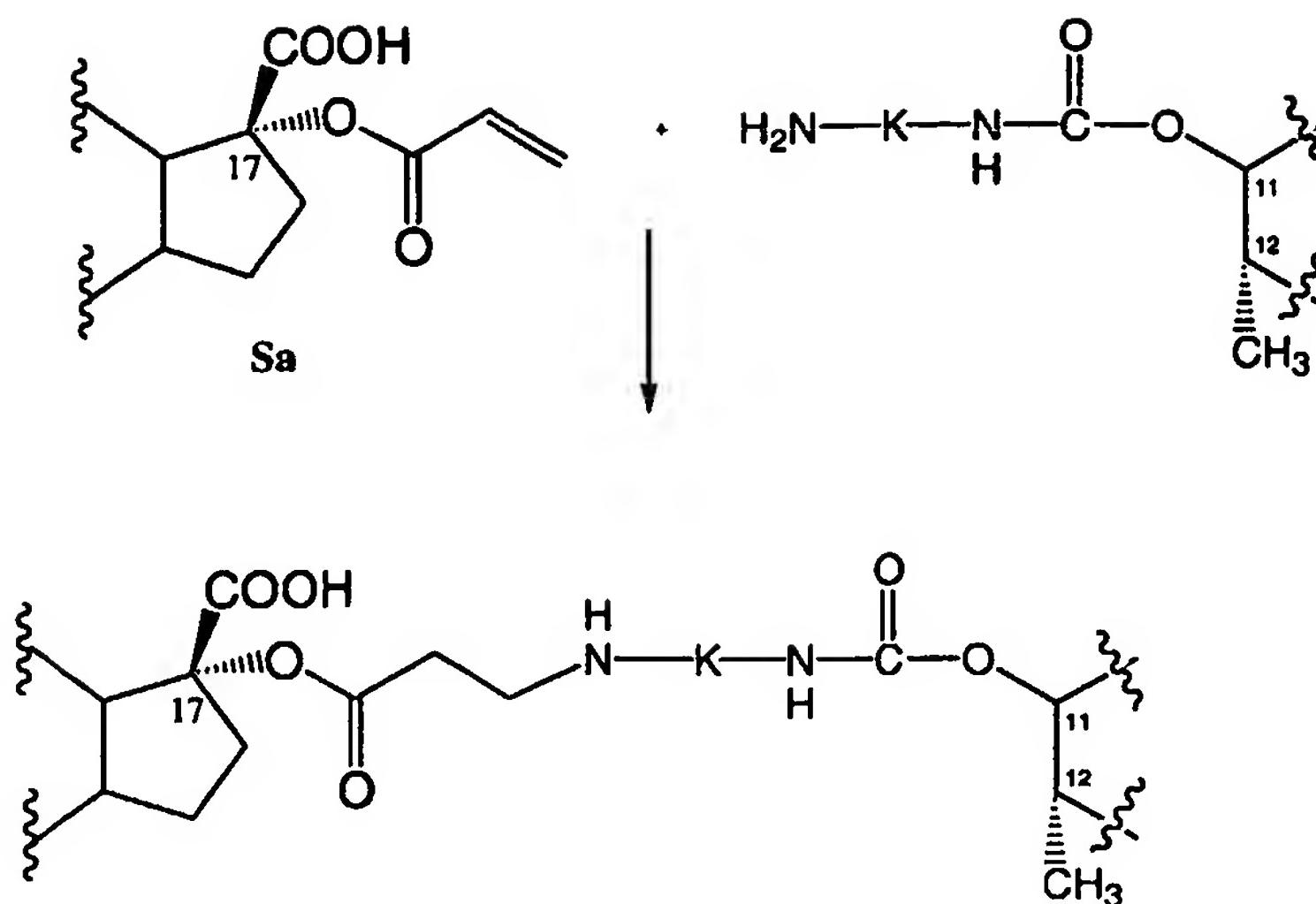
This process may also be performed when the NH group in the macrolide is attached at the 3' position of a sugar ring S¹ (i.e., a desozamine sugar) of the macrolide:



or the 4" position of the sugar ring S²:



Compounds represented by Formula I, where X² is -NH-, can be prepared by reacting a macrolide subunit and a steroid subunit having a -C=C- bond as shown below.



The carboxylic acid group at the 17 position of the starting steroid subunit may be modified prior to the reaction with NH₂-L-M.

The carboxylic acid group at the 17 position of the starting steroid subunit can also be protected prior to the reaction with NH₂-L-M and deprotected after the reaction with NH₂-L-M or the esterification step.

A further aspect of the present invention relates to the methods for using the compounds of Formula I as anti-inflammatory, anti-anaphylactic and immunomodulating agents which can be administered in different ways, depending on the inflammation site. Further, the present invention relates to pharmaceutical compositions containing an effective dose of compounds of the present invention as well as pharmaceutically acceptable excipients, such as carriers or diluents.

The preparation of the pharmaceutical compositions of the invention can include mixing, granulating, tabletting and dissolving the ingredients. Chemical carriers can be in solid or liquid form. Solid carriers can be lactose, sucrose, talc, gelatine, agar, pectin, magnesium stearate, fatty acids without limitation. Liquid carriers can be syrups, oils such as olive, sunflower seed or soybean oils, water, or physiologic saline without limitation. Similarly, carriers may also contain a component for a sustained release of the active component such as glyceryl

monostearate or glyceryl distearate. Several forms of pharmaceutical compositions can be prepared. If a solid carrier is used, these forms can include tablets, caplets, solid gelatinous capsules, powders or granules without limitation that can be administered orally. The amount of the solid carrier can vary but mainly it is in the range from 25 mg to 1 g. If a liquid carrier is used, the formulation can be in the form of a syrup, emulsion, soft gelatinous capsules, or sterile injectable liquids, or nonaqueous liquid suspensions topically or systemically, e.g., orally, parenterally, percutaneously, mucosally, e.g., buccally, intranasally, intrarectally and intravaginally. "Parenterally" means by intravenous, intramuscular or subcutaneous route. The corresponding preparations of the compounds of the present invention can be used in the prophylaxis as well as in the therapeutic treatment (prevention, delay, inhibition or relief) of several disorders (diseases and other pathological inflammatory conditions) caused by or associated with an abnormal or undesirable (excessive, nonregulated, or dysregulated) inflammatory immune response involving the production of inflammatory cytokines or other inflammation mediators, including without limitation TNF- α and IL-1 β . These disorders include autoimmune diseases such as rheumatoid arthritis, insulin-dependent diabetes mellitus, autoimmune thyroiditis, multiple sclerosis, uveoretinitis, lupus erythematosus, scleroderma; other arthritic conditions having an inflammatory component such as rheumatoid spondylitis, osteoarthritis, septic arthritis and polyarthritis; other inflammatory brain disorders, such as meningitis, Alzheimer's disease, AIDS dementia encephalitis, other inflammatory eye inflammations, such as retinitis; inflammatory skin disorders, such as , eczema, other dermatites (e.g., atopic, contact), psoriasis, burns induced by UV radiation (sun rays and similar UV sources); inflammatory bowel disease, such as Crohn's disease, ulcerative colitis; asthma; other allergy disorders, such as allergic rhinitis; conditions associated with acute trauma such as cerebral injury following stroke, heart tissue injury due to myocardial ischemia, lung injury such as that which occurs in adult respiratory distress syndrome; inflammation accompanying infection, such as sepsis, septic shock, toxic shock syndrome, other inflammatory conditions associated with particular organs or tissues , such as nephritis (e.g., glomerulonephritis), inflamed appendix, gout, inflamed gall bladder, chronic obstructive pulmonary disease, congestive heart failure, Type II diabetes, lung fibrosis, vascular disease, such as atherosclerosis and restenosis; and alloimmunity

leading to transplant rejection. The compounds can also be administered by inhalation when application within the respiratory tract is intended. A further object of the present invention relates to the preparation of various pharmaceutical forms of the compounds to achieve the optimal bioavailability of the active compound of **Formula I**.

For percutaneous or mucosal external administration, the compound of **Formula I** can be prepared in a form of an ointment or cream, gel or lotion. Ointments, creams and gels can be formulated using a water or oil base with addition of an appropriate emulsifier or gelling agent. Formulation of the present compounds is especially significant for respiratory inhalation, wherein the compound of **Formula I** is to be delivered in the form of an aerosol under pressure. It is preferred to micronize the compound of **Formula I** after it has been homogenised, e.g., in lactose, glucose, higher fatty acids, sodium salt of dioctylsulfosuccinic acid or, most preferably, in carboxymethyl cellulose, in order to achieve a microparticle size of 5 μm or less for the majority of particles. For the inhalation formulation, the aerosol can be mixed with a gas or a liquid propellant for dispensing the active substance. An inhaler or atomizer or nebulizer may be used. Such devices are known. See, e.g., Newman *et al.*, *Thorax*, 1985, 40:61-676 Berenberg, M., *J. Asthma USA*, 1985, 22:87-92. A Bird nebulizer can also be used. See also U.S. Patents 6,402,733; 6,273,086; and 6,228,346.

The compound of the structure **I** for inhalation is preferably formatted in the form of a dry powder with micronized particles, as described herein.

The compound can also be incorporated into a formulation for treating inflammation localized in an organ or tissue, e.g., Crohn's disease, where it can be administered orally or rectally. Formulations for oral administration can incorporate excipients enabling bioavailability of the compound at the site of inflammation. This can be achieved by different combinations of enteric and delayed release formulations. The compound of **Formula I** can also be used in the treatment of Crohn's disease and intestinal inflammation disease if the compound is applied in the

form of a clyster, for which a suitable formulation can be used, as is well known in the field.

A further aspect of the present invention relates to the use of compounds of Formula I in the treatment of inflammatory diseases, disorders and conditions characterized by or associated with an undesirable inflammatory immune response, especially of all diseases and conditions induced by or associated with an excessive secretion of TNF- α and IL-1.

A therapeutically effective amount of the compound of the present invention can be determined by methods known in the art. Since the compound of the present invention is more efficiently delivered to the desired site than the corresponding anti-inflammatory steroid drug alone, a lesser amount of the compound on a molar basis than of the steroid anti-inflammatory drug can be administered while still achieving the same therapeutic effect. Furthermore, since administration of the compound results in fewer side effects than with the corresponding steroid anti-inflammatory drug, the steroid amount can be increased. Thus, the table below serves only as a guide. A threshold therapeutically effective amount of the compound, a pharmaceutically salt thereof, a solvate thereof, or a prodrug thereof is generally equal to or less than a therapeutically effective amount of the nonsteroidal anti-inflammatory drug on a molar basis. Broad and preferred effective amounts of the compound, a pharmaceutically salt thereof, a solvate thereof, or a prodrug thereof are shown in the table below.

	Amount of Compound, Pharmaceutically Acceptable Salt Thereof, Solvate Thereof, or Prodrug Thereof	
	mg/kg body weight/day of the steroid (had it been administered alone)	μmol/kg body weight/day of the hybrid or the steroid
Broad	from about 0.001 to about 1000	from about 0.004 to about 4000
Preferred	from about 0.01 to about 100	from about 0.04 to about 400
More Preferred	from about 1 to about 100	from about 4 to about 400
Most Preferred	from about 3 to about 30	from about 12 to about 120

For example, if the preferred amount range for prednisone is 1-50 mg/day, this corresponds to a range of 2.79 μmol to 139.5 μmol per day. The starting amount range for a hybrid steroid-macrolide conjugate according to the invention will be also 2.79 μmol to 139.5 μmol of conjugate per day. This dosage can be fine-tuned in light of the present specification using the ordinary skill in the art.

The efficacy of the present compounds can be assessed by any method for assessing inflammation or anti-inflammatory effect. There are many known methods for this purpose including without limitation use of contrast ultrasound in conjunction with injection of microbubbles, measurement of inflammatory cytokines (such as TNF- α , IL-1, IFN- γ) measurement of activated immune system cells (activated T cells, cytotoxic T cells specifically recognizing the inflamed or transplanted tissue) as well as by observation (reduction of oedema reduction of erythema, reduction of pruritus or burning sensation, reduction of body temperature, improvement in function of the afflicted organ) as well as any of the methods provided below as well as any of the methods provided below.

The therapeutic effect of compounds of the present invention was determined in *in vitro* and *in vivo* experiments such as the following.

The beneficial antiinflammatory effect of the compounds of the present invention was determined in the following *in vitro* and *in vivo* experiments:

Assay of Binding to Human Glucocorticoid Receptor

The gene for the alpha isoform of human glucocorticoid receptor (EMBL Acc. No. M10901) was cloned by reverse polymerase chain reaction. The total RNA was isolated from human peripheral blood lymphocytes according to the instructions of the manufacturer (Qiagen), transcribed into cDNA with AMV reverse transcriptase (Roche) and the gene was multiplied by specific primers

- 1) 5'ATATGGATCCCTGATGGACTCCAAAGAACATTAACTCC3' and
- 2) 5'ATATCTCGAGGGCAGTCACTTTGATGAAACAGAAG3'.

The reaction product obtained was cloned into the XhoI/BamHI site of Bluescript KS plasmid (Stratagene), subjected to sequencing by the dideoxy fluorescent method with M13 and M13rev primers (Microsynth) and then it was cloned into the XhoI/BamHI site of pcDNA3.1 hygro(+)plasmid (Invitrogen Life Technologies). 1×10^5 COS-1 cells were seeded onto a 12-well plate (Falcon) in DMEM medium (Invitrogen Life Technologies) with 10% FBS (Biowhitaker) and cultivated to a 70% confluence at 37 °C in an atmosphere with 5% CO₂. The medium was removed and 1 µg of DNA, 7 µl of PLUS reagent and 2 µl of Lipofectamine (Life Technologies) in 500 µl DMEM were added per well. The cells were incubated at 37 °C in an atmosphere with 5% CO₂ and after 5 hours the same volume of 20% FBS/DMEM was added. After 24 hours, the medium was completely changed. 48 hours after transfection, the test compounds in different concentrations and 24 nM [³H]dexamethasone (Pharmacia) in DMEM medium were added. The cells were incubated for 90 minutes at 37 °C in an atmosphere with 5% CO₂, washed three times with PBS buffer (Sigma) cooled to 4 °C (pH=7.4) and then lysed in Tris buffer

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(pH=8.0) (Sigma) with 0.2% SDS (Sigma). After the addition of UltimaGold XR (Packard) scintillation liquid, the residual radioactivity was read in a Tricarb (Packard) β -scintillation counter.

Assay of Inhibition of Mouse T-cell Hybridoma 13 Proliferation as a Result of Apoptosis Induction

In a 96-well plate, triplicates of test steroid dilution in RPMI medium (Instituted of Immunology, Zagreb) with 10% FBS were performed. To the solutions of compounds, 20000 cells per well were added and incubated overnight at 37 °C in an atmosphere with 5% CO₂, then 1 μ Ci of [³H]thymidine (Pharmacia) was added and the mixture was incubated for additional 3 hours. The cells were harvested by applying a vacuum over GF/C filter (Packard). Onto each well, 30 μ l of Microscynt O scintillation liquid (Packard) was added and the incorporated radioactivity was measured on a β -scintillation counter (Packard). The specificity of apoptosis induction by glucocorticoids was proven by antagonising the proliferation inhibition with mifepristone (Sigma).

Balb/c mice are sacrificed by thiopental injection (Pliva Inc.). Spleens are gently sliced and minced on cell strainer. Mononuclear cells are separated by centrifugation using Histopaque 1083 (Sigma diagnostics, Cat No 1083-1) on 400g. Cells are washed with RPMI medium (Institute of immunology, Zagreb, Croatia), and adjusted to 4x10⁶ cells/ml in RPMI medium supplemented with 10% FBS (Invitrogen). Compounds are diluted to appropriate concentrations ranging from 10⁻⁵M to 10⁻¹⁰M in RPMI with 10% FBS, and each dilution (100 μ l) is set in triplicate in 96 well plate. Positive and negative control contained 100 μ l of RPMI with 10% FBS. 50 μ l of murine splenocytes are added on each well, and 50 μ l of concanavalin A (Sigma, cat No C5275), 20 ug/ml in RPMI with 10% FBS in all wells except in negative control where 50 μ l of RPMI with 10% FBS is added.

Plates are placed in incubator (37 °C, 90% relative humidity, 5% CO₂) for 72 hours, and frozen at -70 °C until determination of interleukins.

Interleukins are determined by using sandwich ELISA, using capture and detection antibodies (R&D), according to manufacturer's recommendations.

Model of Lung Eosinophilia in Mice

Male Balb/C mice with a body weight of 20-25 g were randomly divided into groups, and sensitized by an i.p. injection of ovalbumin (OVA, Sigma) on day zero and day fourteen. On the twentieth day, the mice were subjected to a challenge test by i.n. (intranasal) application of OVA (positive control or test groups) or PBS (negative control). 48 hours after i.n. application of OVA, the animals were anaesthetized and the lungs were rinsed with 1 mL of PBS. The cells were separated on Cytospin 3 cytocentrifuge (Shandon). The cells were stained in Diff-Quick (Dade) and the percentage of eosinophils was determined by differential counting of at least 100 cells.

Fluticasone and beclomethasone were used as standard anti-inflammatory substances.

The compounds were administered daily i.n. or i.p. in different doses 2 days before the provocative test and up to the completion of the test. Compounds were administered as suspension either in carboxymethyl cellulose or in lactose solution.

Model of corticosterone suppression and thymus size reduction in rats

Male Wistar rats with a body weight between 200 and 250 g were randomly divided. Tested compounds and standard glucocorticoids were applied by s.c. route once a day for three days. On day three, rats were subjected to cold stress (4 °C, for one hour), anesthetized with Thiopenetal (Pliva Inc.) and blood was taken on heparin. The complete thymus was removed from each animal, and weighed immediately. Plasma was stored at -70 °C until assayed. Corticosterone was extracted with chloroform (5 mL) from 1 mL plasma, or from corticosterone standard dilutions in PBS, interreferent compounds were washed with 0.1 M NaOH, and sulfuric acid:H₂O:C₂H₅OH=8:2:1 was added. Fluorescence was measured 60 minutes later, excitation/emission wavelength was 470/530.

Compounds were considered active if they demonstrated a statistically significant (by Student's t-test, $p<0.05$) result in at least two of the foregoing tests. The molar amounts of compound used were below the threshold amount of macrolide (above $30\mu\text{m}$) for exerting a mild anti-inflammatory effect as reported in the literature.

Uptake/release assay on RAW 264.7, Caco-2 and RBL-2H3 cell line

Cells were seeded in 6 well plate and reached full confluence with 5×10^5 cell per plate. Radiolabelled compounds (app. 50 mCi/mmol each) were loaded on cells in $10 \mu\text{M}$ concentration. Cells were incubated for 2 hours and washed with cold PBS. Cellular concentration was determined immediately by scintillation counting, or were left for another hour with fresh, compound free medium to determine compound release.

SYNTHETIC METHODS AND EXAMPLES

PREPARATION OF INTERMEDIATES

Method A

a) The intermediate **P1** (1.684 g; 2.435 mmole) was dissolved in 50 mL of acrylonitrile and the solution was refluxed at 100°C for 9 hours. Then it was evaporated under reduced pressure and 1.54 g of the raw product **A1** was obtained.

In accordance with the same procedure and starting from the macrolide **P2** (Table 1), nitrile **A2** is obtained. The characteristics of nitrile **A1** and **A2** are given in Table 1.

b) Amine **A3** is obtained starting from the macrolide **A1** (Table 1) by hydrogenation with H₂ with PtO₂.

The macrolide **A1** (1.54 g; 2 mmole) was dissolved in 50 mL of absolute ethanol and hydrated in a reactor with the catalyst PtO₂ (263 mg) under pressure of 40 atm for 24 hours. The reaction mixture was filtered and the solvent was evaporated under reduced pressure. 1.34 g of the mixture was obtained, the mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1 ; 508 mg of amine **A3** was obtained.

In accordance with the same procedure, starting from nitrile **A2**, amine **A4** is obtained.

The characteristics of the macrolides **A3** and **A4** are given in Table 1.

Method B

a) The compound **P3** (1.95 g ; 2.52 mmole) was dissolved in 90 mL of methanol. 1.71 g (12.61 mmole) of NaOAcx3H₂O and 0.68 g (2.64 mmole) of I₂ were added. The reaction mixture was illuminated with a 500W halogen lamp for 2 h. Subsequently, 2-3 drops of 0.1 M Na₂S₂O₃ were added. The solvent was evaporated under reduced pressure and the residue was dissolved in ethyl-acetate and washed with water and with saturated NaHCO₃ solution. The organic layer was dried over anhydrous Na₂SO₄ and evaporated in a rotary evaporator. 3.2 g of the macrolide **B1** was isolated.

b) The compound **B1**, 1.98 g (2.52 mmole), was dissolved in 90 mL of methanol. Subsequently, 3.67 mL (21.09 mmole) of disopropylethylamine and 4.95 mL of ethyl-iodide were added to the solution. The reaction mixture was stirred at 50°C overnight. The solvent was then evaporated under reduced pressure and the residue dissolved in ethyl-acetate and washed with saturated NaHCO₃ solution and water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated. The mixture was purified on a silica gel column in the solvent system CHCl₃ : MeOH : NH₄OH = 6:1:0.1. 559 mg of the compound **B2** was isolated.

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c) In 5 mL of 1,4-diaminobutane, 650 mg (0.82 mmole) of compound **B2** was dissolved. Then, 95 mg (0.82 mmole) of pyridine hydrochloride was added to the solution. The reaction mixture was stirred at room temperature for 3 days. The product was extracted by dichloromethane and washed with water and the organic layer was subsequently dried over Na_2SO_4 and the solvent evaporated under reduced pressure. After purification of the mixture on a silica gel column in the solvent system CHCl_3 : MeOH : $\text{NH}_4\text{OH} = 6:1:0.1$, 300 mg of the amine **B3** was obtained.

According to the same procedure, starting from the macrolide **P3** and diaminobutane, the macrolide **B4** is obtained.

The characteristics of the compounds **B1** to **B4** are given in Table 1.

Method C

a) A suspension of steroid **S1** (110 mg; 0.29 mmole) in dry CH_2Cl_2 (5 mL) cooled to 0°C under argon, was added to 0.380 mL (2.73 mmole) of triethylamine; 80 mg (0.59 mmole) of 1-hydroxybenzotriazole; 230 mg (0.29 mmole) of the amine **A4** and 235 mg (1.23 mmole) of 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride. The reaction mixture was stirred at room temperature in a flow of argon for 24 hours and then evaporated to a smaller volume and purified on a silica gel column (eluent: CHCl_3 : CH_3OH : $\text{NH}_4\text{OH}=6:1:0.1$). 224 mg of the compound **C1** was obtained.

According to a similar procedure, from the macrolide **A4** and the steroid **S2**, the compound **C2** is obtained, from the compound **B4** and the steroid **S2** the compound **C3**, and from the macrolide **A4** and the steroid **S4** the compound **C4**.

The characteristics of compounds **C1** to **C4** are given in Table 1.

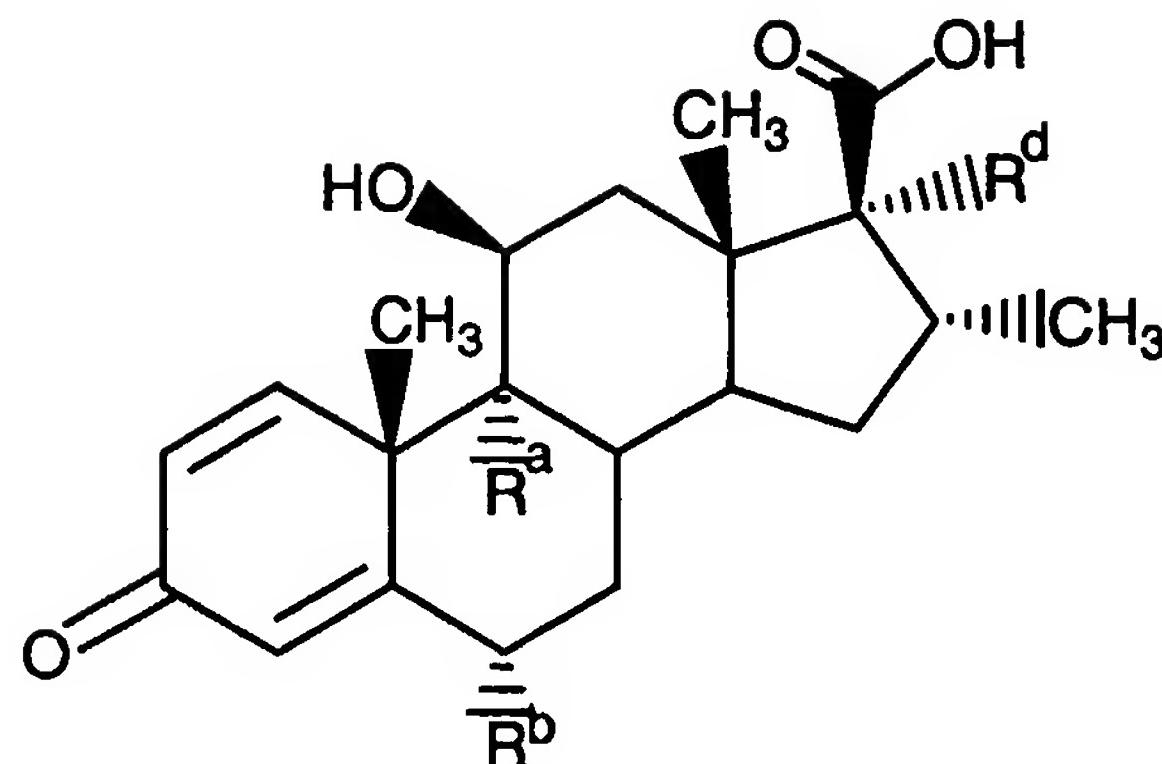
Table 1

Com.	M	R ¹	R ²	R ³	R ⁴	S	Molecular formula	MH ⁺
P1	M1	H	H	H	H	-	C ₃₅ H ₆₅ NO ₁₂	692.9
P2	M2	H	H	H	N(CH ₃) ₂	-	C ₃₇ H ₇₀ N ₂ O ₁₂	735.4
P3	M3	CH ₃	III		N(CH ₃) ₂	-	C ₃₉ H ₇₀ N ₂ O ₁₃	775.5
A1	M1	(CH ₂)CH CN	H	H	H	-	C ₃₈ H ₆₈ N ₂ O ₁₂	745.8
A2	M2	(CH ₂)CH CN	H	H	N(CH ₃) ₂	-	C ₄₀ H ₇₃ O ₁₂ N ₃	789.0
A3	M1	L1	H	H	H	-	C ₃₈ H ₇₂ N ₂ O ₁₂	749.6
A4	M2	L1	H	H	N(CH ₃) ₂	-	C ₄₀ H ₇₇ O ₁₂ N ₃	793.0
B1	M3	CH ₃	III		NHCH ₃	-	C ₃₉ H ₆₉ N ₂ O ₁₃	761.5
B2	M3	CH ₃	III		N(CH ₃)C ₂ H ₅	-	C ₄₀ H ₇₂ N ₂ O ₁₃	789.5
B3	M3	CH ₃	L2	H	N(CH ₃)C ₂ H ₅	--	C ₄₄ H ₉₄ N ₄ O ₁₃	877.6
B4	M3	CH ₃	L2	H	N(CH ₃) ₂	-	C ₄₃ H ₉₂ N ₄ O ₁₃	863.8
C1	M2	L1	H	H	N(CH ₃) ₂	S1	C ₆₁ H ₁₀₂ FN ₃ O ₁₆	1152.6
C2	M2	L1	H	H	N(CH ₃) ₂	S2	C ₆₁ H ₁₀₁ F ₂ N ₃ O ₁₆	1170.6
C3	M3	CH ₃	L2	H	N(CH ₃) ₂	S2	C ₆₄ H ₁₀₆ F ₂ N ₄ O ₁₇	1241.7
C4	M2	L1	H	H	N(CH ₃) ₂	S4	C ₆₁ H ₁₀₂ FN ₃ O ₁₅	1136.7

L1 = -(CH₂)₃-NH₂

L2 = -C(O) NH-(CH₂)₄-NH₂

M1-M3 corresponds to the macrolide subunit of the formula immediately preceding the foregoing table wherein the variables R1-R4 are as specified in Table 1.

Table 2

S	R ^a	R ^b	R ^d	Molecular formula	MH ⁺	MH ⁻
S1	F	H	OH	C ₂₁ H ₂₇ FO ₅	378.9	-
S2	F	F	OH	C ₂₁ H ₂₆ O ₅ F ₂	397.4	395.3
S3	Cl	F	OH	C ₂₁ H ₂₆ O ₅ ClF	-	411.3
S4	F	H	H	C ₂₁ H ₂₇ FO ₄	-	361.2

Example 1

Compound 1: I; M = M1, L = L1, S = S1; (R² = R³ = R⁴ = H, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

In 5 mL of dry dichlormethane, 76 mg of the steroid **S1** (0.2 mmole) was dissolved in an inert atmosphere. Subsequently, 0.25 mL of triethylamine, 53 mg of hydroxybenzotriazole, 150 mg of the macrolide **A3** (0.2 mmole) and 157 mg of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride was added to the solution. The reaction mixture was stirred at room temperature overnight. The solvent was

evaporated under reduced pressure. The mixture obtained was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 167 mg of the compound 1 was obtained; MS (*m/z*): 1109.7 [MH]⁺. IR(cm⁻¹)/KBr: 3425, 2974, 2939, 2875, 1721, 1665, 1627, 1525, 1459, 1379, 1296, 1262, 1168, 1125, 1064, 1035, 1013, 959, 927, 894, 804.

Example 2

Compound 2: I; M =M1, L = L1, S =S2; (R² = R³ = R⁴ = H, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = R^b = F, R^d = OH)

In 5 mL of dry dichlormethane, 80 mg of the steroid S2 (0.2 mmole) was dissolved in an inert atmosphere. Subsequently, 0.25 mL of triethylamine, 53 mg of hydroxybenzotriazole, 150 mg of the macrolide A3 (0.2 mmole) and 157 mg of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride was added to the solution. The reaction mixture was stirred at room temperature overnight. The solvent was evaporated under reduced pressure and the mixture obtained was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 129 mg of the compound 2 was obtained; MS (*m/z*): 1127.7 [MH]⁺. IR(cm⁻¹)/KBr: 3424, 2972, 2939, 2876, 1719, 1670, 1631, 1560, 1523, 1458, 1379, 1317, 1264, 1167, 1127, 1064, 1033, 997, 960, 900, 820, 735, 709.

Example 3

Compound 3: I; M = M1, L = L1, S = S3; (R² = R³ = R⁴ = H, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = Cl, R^b = F, R^d = OH)

80 mg of the steroid S3 (0.19 mmole) was dissolved in 5 mL of dry dichlormethane in an inert atmosphere. 0.25 mL of triethylamine, was added to the solution, which became clear. Subsequently, 53.5 mg of hydroxybenzotriazole, 145 mg of the macrolide A3 (0.19 mmole) and 157 mg of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride were added. The reaction mixture was stirred at room temperature overnight. The solvent was evaporated under reduced pressure and the obtained mixture was purified on a silica gel column, eluent

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$\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH}=6:1:0.1$. 200 mg of the compound **3** was obtained; MS (*m/z*): 1143.6 [MH]⁺. IR(cm^{-1})/KBr: 3423, 2971, 2937, 2876, 1719, 1668, 1630, 1561, 1523, 1459, 1379, 1319, 1262, 1168, 1125, 1064, 998, 959, 901, 819, 756.

Example 4

Compound 4: I; M = M3, L = L2, S = S1; (R¹ = CH₃, R³ = H, R⁴ = N(CH₃)CH₂CH₃, X¹ = -C(O)NH-, m = 4, X² = -NH-, R^a = F, R^b = H, R^d = OH).

In 15 mL of dichloromethane, 126 mg (0.33 mmole) of the compound **S1** was dissolved. To the reaction mixture were added, 0.363 mL of triethylamine, 90 mg of hydroxybenzotriazole, 292 mg (0.33 mmole) of the compound **B3** and 229 mg of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride. The reaction mixture was stirred at room temperature in an inert atmosphere for 24 h. The solvent was then evaporated, and the compound was purified on a silica gel column in the solvent system $\text{CHCl}_3 : \text{MeOH} : \text{NH}_4\text{OH} = 6:1:0.1$. 58 mg of the compound **4** was isolated: MS (*m/z*): 1237.8 [MH]⁺. IR(cm^{-1})/KBr: 3437, 2973, 2938, 1723, 1658, 1545, 1526, 1462, 1379, 1271, 1172, 1111, 1053, 996, 895, 807.

Example 5

Compound 5: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = NHCH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

Procedure 1:

In 3 mL of methanol, 100 mg of the compound **C1** (0.086 mmole) was dissolved. To the solution, 59 mg of sodiumacetate trihydrate and 23 mg of iodine was added. The reaction mixture was illuminated with a 500W halogen lamp for 2 hours. Subsequently, a couple of drops of 1M sodium-thiosulphate were added. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column with, eluent $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH}=6:1:0.1$. 21 mg of the compound **5** was obtained; MS (*m/z*): 1138,5 [MH]⁺. IR(cm^{-1})/KBr: 3423, 2972,

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2938, 2875, 1719, 1664, 1625, 1560, 1541, 1523, 1459, 1379, 1243, 1169,
1125, 1072, 1012, 959, 894, 807, 755.

Procedure 2:

The compound **C1** (500 mg; 0.43 mmole) was dissolved in 30 mL of acetonitrile. Subsequently, 146 mg of N-iodine succinimide (0.65 mmole) at 0°C and in a flow of argon was added to the solution. The reaction mixture was heated to room temperature and stirred for 5 hours. Subsequently, the mixture was diluted with 100 mL of dichlormethane and extracted with 100 mL of 1:1 5% NaHSO₃ : Na₂CO₃ and saturated NaCl solution. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated in a rotary evaporator. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 155 mg of the compound **5** was obtained; MS (*m/z*): 1139.3 [MH]⁺.

Example 6

Compound 6: I; M = M2, L = L1, S = S2; (R² = R³ = H, R⁴ = NHCH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = R^b = F, R^d = OH)

The compound **C2** (250 mg; 0.21 mmole) was dissolved in 10 mL of methanol. To the solution was added, 145 mg of sodiumacetate trihydrate (1 mmole) and 57.4 mg of iodine (0.22 mmole). The reaction mixture was illuminated with a 500W halogen lamp for 2 hours. Several drops of 1M sodium-thiosulphate were added in order to destroy the surplus iodine. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH= 6:1:0.1. 83 mg of the compound **6** was obtained; MS (*m/z*): 1156.7 [MH]⁺. IR(cm⁻¹)/KBr: 3432, 2972, 2938, 2876, 1774, 1719, 1670, 1655, 1578, 1560, 1523, 1459, 1378, 1317, 1263, 1168, 1127, 1070, 1034, 998, 959, 899, 820, 755, 710, 669.

Example 7

Compound 7: I; M = M3, L = L2, S = S2; (R¹ = CH₃, R³ = H, R⁴ = NHCH₃,

X¹ = -C(O)NH-, m = 4, X² = -NH-, R^a = R^b = F, R^d = OH)

The compound C3 (74 mg; 0.06 mmole) was dissolved in 10 mL of methanol. 38 mg (0.28 mmole) of NaOACx3H₂O and 15 mg (0.06 mmole) of I₂ were added. The reaction mixture was illuminated with a 500W halogen lamp for 2 h. Then, 2-3 drops of 0.1 M Na₂S₂O₃ were added. The solvent was then evaporated under reduced pressure and the residue was dissolved in ethyl-acetate and washed with water and saturated NaHCO₃ solution. The organic layer was dried over anhydrous Na₂SO₄ and evaporated. The product was purified on a silica gel column in the solvent system CHCl₃ : MeOH : NH₄OH = 6:1:0.1. The quantity of 60 mg of the compound 7 was isolated; MS (m/z): 1226.5 [MH]⁺. IR(cm⁻¹)/KBr: 3424, 2972, 2939, 2876, 1719, 1670, 1631, 1560, 1523, 1459, 1379, 1314, 1259, 1167, 1127, 1068, 1002, 947, 899, 821, 710.

Example 8

Compound 8: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = N(CH₃)CH₂CH₃,

X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

The compound 5 (100 mg; 0.09 mmole) was dissolved in 3mL of methanol. To the solution, 127 µl of N,N-diisopropylethylamine and 45 µl of ethyliodide was added. The reaction mixture was stirred at a temperature of 50°C for 20 hours. Subsequently, the mixture was diluted with 30 mL of ethyl-acetate and washed with 30 mL of saturated aqueous sodium-hydrogencarbonate solution and 30 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated by reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH =6:1:0.1. 45 mg of the compound 8 was obtained; MS (m/z): 1166.5 [MH]⁺. IR(cm⁻¹)/KBr: 3411, 3238, 2969, 2936, 2873, 1723, 1664, 1625, 1565, 1528, 1462, 1379, 1327, 1259, 1169, 1063, 1012, 959, 929, 894, 820, 749, 663, 617.

Example 9

Compound 9: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = N(CH₃)CH₂CH₂CH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

The compound 5 (90 mg; 0.08 mmole) was dissolved in 2 mL of methanol. To the solution, 115 µl of N,N-diisopropylethylamine (0.67 mmole) and 45.7 µl of propylbromide (0.50 mmole) was added. The reaction mixture was stirred at a temperature of 50°C for 20 hours. Subsequently, the mixture was diluted with 20 mL of ethyl-acetate and washed with 20 mL of saturated aqueous sodium-hydrogencarbonate solution and 20 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. The quantity of 20 mg of the compound 9 was obtained; MS (m/z): 1180.6 [MH]⁺. IR(cm⁻¹)/KBr: 3432, 2964, 2927, 2881, 1736, 1666, 1639, 1627, 1562, 1545, 1525, 1460, 1379, 1262, 1168, 1099, 1054, 1017, 893, 802, 701.

Example 10

Compound 10: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = N(CH₃)CH₂CH₂CH₂CH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

In 3mL of methanol, 100 mg of the compound 5 (0.09 mmole) was dissolved. To the solution, 128 µl of N,N-diisopropylethylamine (0.75 mmole) and 63.5 µl of butyl-iodide (0.56 mmole) was added. The reaction mixture was stirred at a temperature of 50°C for 20 hours. It was then diluted with 25 mL of ethyl-acetate and washed with 25 mL of saturated aqueous sodium-hydrogencarbonate solution and 25 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated by reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 50 mg of compound 10 was obtained; MS (m/z): 1195.1 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2937, 2874, 1774, 1735,

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 1719, 1664, 1638, 1578, 1560, 1523, 1491, 1459, 1378, 1263, 1244, 1169,
 1106, 1055, 1013, 959, 894, 805, 754.

Example 11

Compound 11: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = N(CH₃)CH(CH₃)₂, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

The compound 5 (81 mg; 0.07 mmole) was dissolved in 3 mL of methanol. To the solution, 26 µl of acetone, 6.8 mg of sodium-cyanoborhydride and a drop of acetic acid was added. The reaction mixture was stirred at room temperature for 2 days. After two days, an additional 26 µl of acetone was added. The reaction mixture was stirred for further 3 days. The reaction mixture was extracted with 20 mL of ethyl-acetate and 20 mL of 50% aqueous sodium-hydrogencarbonate solution. The organic layer was washed with water and with saturated aqueous NaCl solution and dried over anhydrous sodium-sulphate. The solvent was evaporated in a rotary evaporator. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 11 mg of the compound 11 was obtained; MS (m/z): 1180.9 [MH]⁺.

Example 12

Compound 12: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = cyclohexyl, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

The compound 5 (83 mg; 0.07 mmole) was dissolved in 2 mL of methanol. To the solution, 38 µl of cyclohexanone (0.36 mmole), 7 mg of sodium-cyanoborhydride (0.11 mmole) and a drop of acetic acid was added. The reaction mixture was stirred at room temperature for 2 days. The reaction mixture was extracted with 40 mL of ethyl-acetate and 40 mL of 50% aqueous sodium-hydrogencarbonate solution. The organic layer was washed with water and saturated aqueous NaCl solution and dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent

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 $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH}=6:1:0.1$. 11 mg of the compound **12** was obtained; MS (*m/z*): 1220.6 [MH]⁺.

Example 13

Compound 13: I; M = M2, L = L1, S = S2; (R² = R³ = H, R⁴ = benzyl, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = R^b = F, R^d = OH)

The compound **6** (85 mg; 0.07 mmole) was dissolved in 2 mL of methanol. To the solution, 107 µl of N,N-diisopropylethylamine (0.62 mmole) and 55.5 µl of benzylbromide (0.47 mmole) was added. The reaction mixture was stirred at a temperature of 50°C for 20 hours and diluted with 20 mL of ethyl-acetate and washed with 20 mL of saturated aqueous sodium-hydrogencarbonate solution and 20 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH}=6:1:0.1$. 18 mg of compound **13** was obtained; MS (*m/z*): 1246.6 [MH]⁺. IR(cm⁻¹)/KBr: 3433, 2963, 2934, 2875, 1729, 1668, 1640, 1564, 1528, 1496, 1456, 1378, 1318, 1261, 1167, 1126, 1108, 1053, 1034, 999, 960, 901, 802, 750, 700.

Example 14

Compound 14: I; M = M2, L = L1, S = S1; (R² = R³ = H, R⁴ = N(CH₃)COCH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

In 5 mL of methanol, 87 mg of the compound **5** was dissolved and cooled to 2°C. At this temperature, 20 µl of acetanhydride was added drop wise to the reaction mixture. After stirring for three hours at this temperature, the solvent was evaporated and a white, oily product was obtained which was subsequently purified on a silica gel column, eluent $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH}=6:1:0.1$. 69 mg of compound **14** was obtained; MS (*m/z*): 1180.5 [MH]⁺. IR(cm⁻¹)/KBr: 3422, 2970, 2938, 2875, 1794, 1774, 1710, 1686, 1664, 1625, 1560, 1523, 1458, 1364, 1223, 1168, 1124, 1061, 1012, 959, 895, 669.

Example 15

Compound 15: I; M = M3, L = L2, S = S2; (R¹ = CH₃, R³ = H, R⁴ = N(CH₃)CH₂CH₃, X¹ = -C(O)NH-, m = 4, X² = -NH-, R^a = R^b = F, R^d = OH)

The compound 7 (60 mg; 0.05 mmole) was dissolved in 3 mL of methanol. Subsequently, 0.100 mL (0.57 mmole) of diisopropylethylamine and 0.233 mL of ethyl-iodide was added to the solution. The reaction mixture was stirred at 50°C overnight. The solvent was then evaporated under reduced pressure and the residue was dissolved in ethyl-acetate and washed with saturated NaHCO₃ solution and water. The organic layer was dried over anhydrous Na₂SO₄ and evaporated. The mixture was purified on a silica gel column in the solvent system CHCl₃ : MeOH : NH₄OH = 6:1:0.1. 15 mg of compound 15 was isolated; MS (m/z):1255.8 [MH]⁺.

Example 16

Compound 16: I; M = M2, L = L1, S = S4; (R² = R³ = H, R⁴ = NHCH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = H)

The compound C4 (590 mg; 0.519 mmole) was dissolved in 50 mL of acetonitrile. At a temperature of 0°C and in a flow of argon, 175 mg of N-iodosuccinimide (0.78 mmole) was added to the solution. The reaction mixture was heated to room temperature and stirred for 5 hours. Subsequently, the mixture was diluted with 100 mL of dichlormethane and extracted with 100 mL of 1:1 5% NaHSO₃ : Na₂CO₃ and saturated NaCl solution. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH= 6:1:0.1. 100 mg of compound 16 was obtained; MS (m/z): 1123.2 [MH]⁺. IR(cm⁻¹)/KBr: 3448, 2969, 2939, 2876, 1719, 1664, 1638, 1629, 1560, 1542, 1509, 1500, 1459, 1379, 1294, 1247, 1168, 1124, 1065, 1011, 959, 891, 829, 808, 755, 670.

Example 17

Compound 17: I; M = M2, L = L1, S = S4; (R² = R³ = H, R⁴ = N(CH₃)CH₂CH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = H)

The compound **16** (70 mg; 0.06 mmole) was dissolved in 2 mL of methanol. To the solution was added, 90.8 µl of N,N-diisopropylethylamine (0.53 mmole) and 119.4 µl of ethyl iodide (0.39 mmole). The reaction mixture was stirred in a magnetic mixer at a temperature 50°C for 20 hours. Subsequently, the mixture was diluted with 30 mL of ethyl-acetate and washed with 30 mL of saturated aqueous sodium-hydrogen carbonate solution and 30 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 32 mg of compound **17** was obtained; MS (*m/z*): 1150.8 [MH]⁺. IR(cm⁻¹)/KBr: 3434, 2969, 2939, 2870, 1719, 1664, 1630, 1561, 1535, 1458, 1383, 1327, 1293, 1236, 1168, 1109, 1058, 1012, 959, 890, 812, 731, 704.

Example 18

Compound 18: I; M = M2, L = L1, S = S4; (R² = R³ = H, R⁴ = N(CH₃)COCH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = H)

The compound **16** (120 mg) was dissolved in 5 mL of methanol and cooled to 2°C. At this temperature, into the reaction mixture, 28 µl of acetanhydride (0.2 mmole) was added dropwise. After stirring for three hours at this temperature, the solvent was evaporated and the residue was recrystallized (chloroform/hexane). 148 mg of compound **18** was obtained; MS (*m/z*): 1180.5 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2979, 2938, 2870, 1735, 1719, 1664, 1625, 1560, 1542, 1509, 1459, 1381, 1293, 1248, 1168, 1122, 1060, 1012, 891, 830, 647.

Example 19

Compound 19: I; M = M2, L = L1, S = S2; (R² = R³ = H, R⁴ = N(CH₃)CH₂CH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = R^b = F, R^d = OH)

The compound **6** (68 mg; 0.06 mmole) was dissolved in 2 mL of methanol. To the solution was added, 85 µl of N,N-diisopropylethylamine (0.5 mmole) and 112 µl of ethyliodide (0.4 mmole). The reaction mixture was stirred at a temperature of 50°C for 20 hours. It was then diluted with 20 mL of ethyl-acetate and washed with 20 mL of saturated aqueous sodium-hydrogencarbonate solution and 20 mL of water. The organic layer was dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH=6:1:0.1. 38 mg of compound **19** was obtained; MS (m/z): 1185.7 [MH]⁺. IR(cm⁻¹)/KBr: 3425, 2964, 2870, 1721, 1671, 1638, 1562, 1544, 1525, 1460, 1379, 1262, 1168, 1092, 1055, 1032, 958, 899, 864, 803, 707.

Example 20

Compound 20: I; M = M2, L = L1, S = S1; (R² and R³ = III, R⁴ = N(CH₃)₂, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

Compound **C1** (300 mg; 0.26 mmole) was mixed with 139 mg of ethylencarbonate (1.58 mmole) and 42.7 mg of potassium-carbonate (0.31 mmole). To the reaction mixture, 5 mL of ethyl-acetate was added. The solution was heated to 75°C under stirring for 2 days. Crude potassium-carbonate was filtered, the filtrate was diluted with ethyl-acetate and extracted with water. The organic layer was dried over anhydrous sodium sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column (solvent system: CHCl₃:MeOH:NH₄OH= 6:1:0.1) and 72 mg of compound **20** was obtained; MS (m/z): 1178.3 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2973, 2939, 2874, 1806, 1736, 1685, 1665, 1625, 1560, 1523, 1458, 1380, 1240, 1168, 1110, 1075, 1052, 1014, 893, 834, 755, 687.

Example 21

Compound 21: I; M = M2, L = L1, S = S1; (R² and R³ = III, R⁴ = N(CH₃)CH₂CH₃, X¹ = -CH₂-, m = 2, X² = -NH-, R^a = F, R^b = H, R^d = OH)

Compound 8 (65 mg; 0.06 mmole) was mixed with 30 mg of ethylenecarbonate (0.34 mmole) and 9.2 mg of potassium-carbonate (0.06 mmole). To the reaction mixture, 3 mL of ethyl-acetate was added. The solution was heated to 75°C under stirring for 3 days. Crude potassium-carbonate was filtered, the filtrate was diluted with ethyl-acetate and extracted with water. The organic layer was dried over anhydrous sodium sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH = 6:1:0.1. 10 mg of compound 21 was obtained; MS (m/z): 1192.9 [MH]⁺.

Example 22

Compound 22: I; M = M3, L = L2, S = S4; (R¹ = CH₃, R³ = H, R⁴ = N(CH₃)CH₂CH₃, X¹ = -C(O)NH-, m = 4, X² = -NH-, R^a = F, R^b = H, R^d = H)

In 10 mL of dichloromethane, 0.07 g (0.21 mmole) of the compound S4 was dissolved. Subsequently, 0.226 mL (1.62 mmole) of triethylamine, 56 mg (0.42 mmole) of hydroxybenzotriazole, 182 mg (0.21 mmole) of the compound B3 and 143 mg (0.83 mmole) of 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride was added to the reaction mixture. The reaction mixture was stirred at room temperature in an inert atmosphere for 24 h. The solvent was then evaporated and the compound purified on a silica gel column in the solvent system CHCl₃ : MeOH : NH₄OH = 6:1:0.1. 52 mg of compound 22 was isolated; MS (m/z): 1221.8 [MH]⁺. IR(cm⁻¹)/KBr: 3450, 2971, 2938, 2875, 1775, 1721, 1705, 1628, 1562, 1544, 1526, 1460, 1378, 1294, 1254, 1167, 1110, 1054, 1013, 889, 815, 755.

PREPARATION PROCESSES WITH EXAMPLES

PREPARATION OF INTERMEDIATES

Method AA

The compound **P1A** (2 g; 2.46 mmole) was dissolved in 100 mL of methanol. To the solution, 2.2 g of sodium acetate trihydrate and 900 mg of iodine was added. The reaction mixture was illuminated with a 500 W halogen lamp for 2 hours. Then, a couple of drops of 1M sodium thiosulphate were added. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column (eluent: $\text{CHCl}_3:\text{CH}_3\text{OH}:\text{NH}_4\text{OH} = 6:1:0.1$). The quantity of 870 mg of the compound **A1A** was obtained.

According to the above procedure, starting from the macrolide **P2A** (Table 3), the compound **A2A** is obtained.

The characteristics of the compounds **A1** and **A2** are given in Table 3.

Method BB

The compound **A1** (4.9 g; 6.68 mmole) was dissolved in 100 mL of methanol. To the solution, 36 mL of N,N-diisopropylethylamine and 13 mL of ethyl iodide were added. The reaction mixture was stirred at the temperature of 50°C for 20 hours. The reaction mixture was then diluted with ethyl acetate and washed with saturated aqueous solution of sodium-hydrogen carbonate and water. The organic extract was dried over anhydrous sodium sulphate. The solvent was evaporated under reduced pressure. The obtained mixture was purified on a silica gel column (eluent: $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH} = 6:1:0.1$). The quantity of 1.63 g of the compound **B1A** was obtained.

According to the above procedure, starting from the compound **A2A**, the compound **B2A** is obtained.

The compound **A1A** (770 mg; 1.048 mmole) was dissolved in 30 mL of acrylonitrile and the reaction mixture was refluxed for 10 hours. Subsequently, acrylonitrile was evaporated under reduced pressure. 827 mg of compound **B3A** was obtained.

In accordance with the above procedure and starting from the compound **A2**, the compound **B4A** is obtained.

The characteristics of compounds **B1A** to **B4A** are given in Table 1.

Method CC

In a 100 mL flask, the macrolide **P1A** (6.4 g; 8.6 mmole) was dissolved in 50 mL of ethyl acetate. Acetanhydride (1.22 mL; 13.0 mmole) was then added to the reaction mixture. The mixture was stirred at room temperature for 18 h. The reaction mixture was diluted with 50 mL of water and stirred for additional 30 minutes. The pH of the aqueous layer was adjusted to 2.5. Organic and aqueous layers were separated and then the pH of the aqueous phase was adjusted to 9.5 and the aqueous layer was then extracted with ethyl acetate. The extracts were dried over anhydrous Na₂SO₄. The solvent was evaporated under reduced pressure and 3.33 g of compound **C1A** was obtained.

According to the above procedure and starting individually from compounds **P2A**, **P3A**, **B1A** or **B2A**, the compounds **C2A**, **C3A**, **C4A** or **C5A** are obtained.

The characteristics of the compounds **C1A** to **C5A** are given in Table 1.

Method DD

The compound **C1A** (6.8 g; 8.6 mmole) was dissolved in 70 mL of dichloromethane. Added to the mixture was 6.8 mL of DMSO and 1-(3-dimethyl aminopropyl)-3-ethyl-carbodiimide hydrochloride (5 g; 29 mmole) in an inert atmosphere. Pyridine trifluoroacetate, dissolved in 3 mL of dichloromethane, was then gradually added drop wise. The reaction mixture was stirred for 2 h at room

temperature and subsequently water was added into the reaction mixture. The layers were separated, and the pH of the aqueous layer was adjusted to 4.0. After extraction with ethyl acetate, the pH was adjusted to 6.5. Extraction was repeated, and the pH of the aqueous layer was adjusted to 9.5. After extraction with ethyl acetate, the organic layer was dried over anhydrous Na_2SO_4 . The solvent was evaporated and 2 g of compound **D1A** was isolated.

According to the above procedure and starting individually from the compounds **C2A**, **C3A**, **C4A** or **C5A**, the compounds **D2A**, **D3A**, **D4A**, or **D5A** are obtained.

The characteristics of the compounds **D1A** to **D5A** are given in Table 1.

Method EE

The solution of the compound **D1A** (1.5 g ;1.9 mmole) in 82 mL of methanol was stirred at room temperature for 20 h. The solvent was then evaporated under reduced pressure and 1.2 g of the compound **E1A** was obtained.

According to the above procedure and starting individually from the compounds **D2A**, **D3A**, **D4A** or **D5A**, the compounds **E2A**, **E3A**, **E4A** or **E5A** are obtained.

The characteristics of the compounds **E1A** to **E5A** are given in Table 1.

Method FF

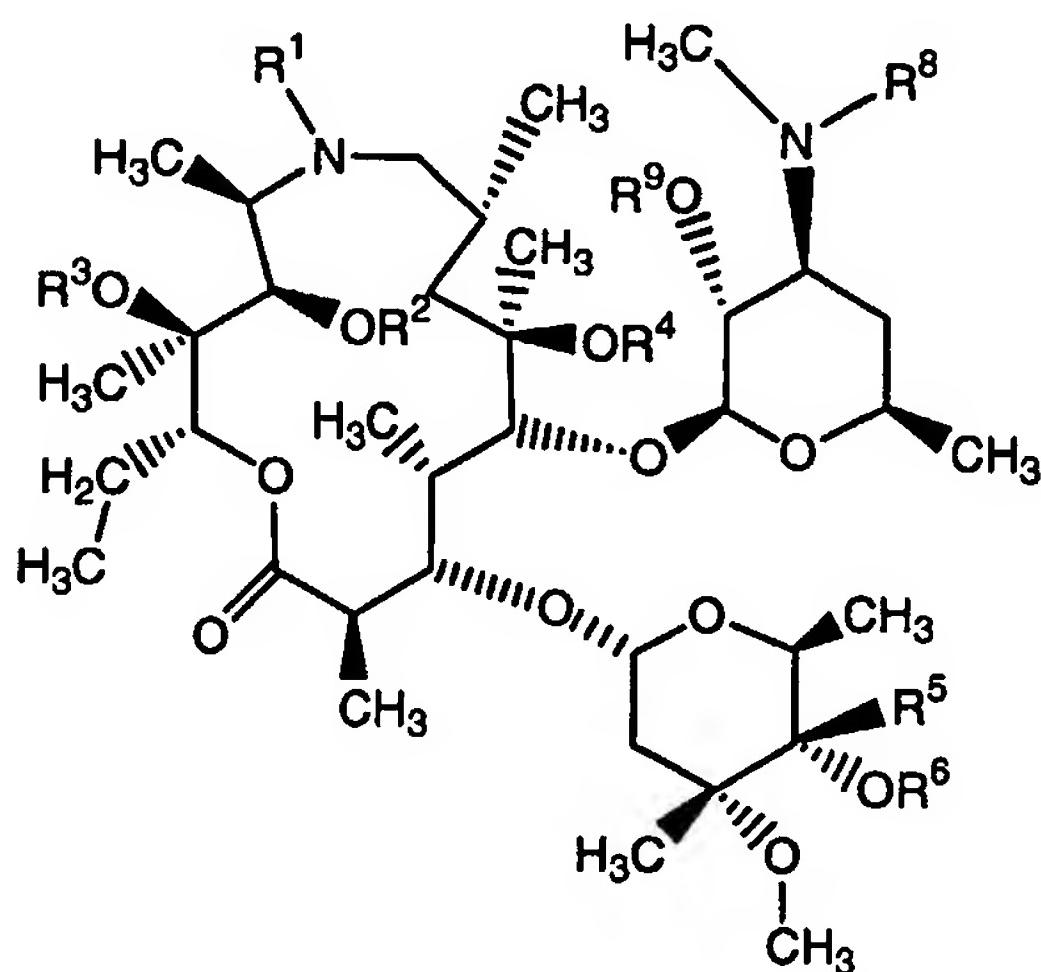
To the solution of NaH (30.9 mg) dissolved in 3.7 mL of DMF, 297 mg of trimethylsulphonium iodide (TMSI) was added. After 15 minutes, the solution of the compound **E1A** (700 mg; 0.94 mmole), dissolved in 2.3 mL of DMSO, was slowly added drop wise. The mixture was stirred at room temperature for 15 min, and then at 55°C for 45 min. Stirring was then continued at room temperature overnight. The reaction mixture was poured into the mixture of water and ethyl acetate. The organic layer was separated, washed with water and saturated solution of NaCl, dried

over Na_2SO_4 , and the solvent was then evaporated under reduced pressure. The quantity of 0.53 g of the compound **F1A** was obtained.

According to the above procedure, starting individually from the compounds **E2A**, **E3A**, **E4A** or **E5A**, the compounds **F2A**, **F3A**, **F4A** or **F5A** are obtained.

The characteristics of the compounds **F1A** to **F5A** are given in Table 3.

Table 3



Com.	R ¹	R ²	R ³	R ⁴	R ⁵	R ⁶	R ⁸	R ⁹	Molecular formula	MH ⁺
P1A	CH ₃	H	H	H	H	H	CH ₃	H	C ₃₉ H ₇₂ N ₂ O ₁₂	749.5
P2A	H	C=O		CH ₃	H	H	CH ₃	H	C ₃₈ H ₇₀ N ₂ O ₁₃	763.5
P3A	IPA ^a	H	H	H	H	H	CH ₃	H	C ₄₁ H ₇₇ N ₃ O ₁₃	820.5
A1A	CH ₃	H	H	H	H	H	H	H	C ₃₇ H ₇₀ N ₂ O ₁₂	735.5

A2A	H	C=O		CH ₃	H	H	H	H	C ₃₇ H ₆₉ N ₂ O ₁₃	749.4
B1A	CH ₃	H	H	H	H	H	C ₂ H ₅	H	C ₃₉ H ₇₄ N ₂ O ₁₂	763.6
B2A	H	C=O		CH ₃	H	H	C ₂ H ₅	H	C ₃₉ H ₇₂ N ₂ O ₁₃	777.7
B3A	CH ₃	H	H	H	H	H	(CH ₂) ₂ -CN	H	C ₄₀ H ₇₃ N ₃ O ₁₂	788.5
B4A	H	C=O		CH ₃	H	H	(CH ₂) ₂ -CN	H	C ₄₀ H ₇₁ N ₃ O ₁₃	822.8
C1A	CH ₃	H	H	H	H	H	CH ₃	Ac	C ₄₀ H ₇₄ N ₂ O ₁₃	792.1
C2A	H	C=O		CH ₃	H	H	CH ₃	Ac	C ₄₀ H ₇₂ N ₂ O ₁₄	805.5
C3A	IPA	H	H	H	H	H	CH ₃	Ac	C ₄₃ H ₇₉ N ₃ O ₁₄	862.6
C4A	CH ₃	H	H	H	H	H	C ₂ H ₅	Ac	C ₄₁ H ₇₆ N ₂ O ₁₃	805.6
C5A	H	C=O		CH ₃	H	H	C ₂ H ₅	Ac	C ₄₁ H ₇₄ N ₂ O ₁₄	819.6
D1A	CH ₃	H	H	H	C=O		CH ₃	Ac	C ₄₀ H ₇₂ N ₂ O ₁₃	789.6
D2A	H	C=O		CH ₃	C=O		CH ₃	Ac	C ₄₀ H ₇₀ N ₂ O ₁₄	803.8
D3A	IPA	H	H	H	C=O		CH ₃	Ac	C ₄₃ H ₇₇ N ₃ O ₁₄	860.8
D4A	CH ₃	H	H	H	C=O		C ₂ H ₅	Ac	C ₄₁ H ₇₄ N ₂ O ₁₃	803.6
D5A	H	C=O		CH ₃	C=O		C ₂ H ₅	Ac	C ₄₁ H ₇₂ N ₂ O ₁₄	817.6
E1A	CH ₃	H	H	H	C=O		CH ₃	H	C ₃₉ H ₇₀ N ₂ O ₁₂	747.6

E2A	H	C=O	CH ₃	C=O	CH ₃	H	C ₃₉ H ₆₉ N ₂ O ₁₃	761.5	
E3A	IPA	H	H	H	C=O	CH ₃	H	C ₄₁ H ₇₅ N ₃ O ₁₃	818.5
E4A	CH ₃	H	H	H	C=O	C ₂ H ₅	H	C ₃₉ H ₇₂ N ₂ O ₁₂	761.6
E5A	H	C=O	CH ₃	C=O	C ₂ H ₅	H	C ₃₉ H ₇₀ N ₂ O ₁₃	775.5	
F1A	CH ₃	H	H	H	EPO ^b	CH ₃	H	C ₃₉ H ₇₂ N ₂ O ₁₂	761.8
F2A	H	C=O	CH ₃	EPO	CH ₃	H	C ₃₉ H ₇₀ N ₂ O ₁₃	775.5	
F3A	IPA	H	H	H	EPO	CH ₃	H	C ₄₂ H ₇₇ N ₃ O ₁₃	832.5
F4A	CH ₃	H	H	H	EPO	C ₂ H ₅	H	C ₄₀ H ₇₄ N ₂ O ₁₂	775.6
F5A	H	C=O	CH ₃	EPO	C ₂ H ₅	H	C ₄₀ H ₇₂ N ₂ O ₁₃	789.5	

a) IPA = C(O)NHCH(CH₃)₂

b) EPO = Oksiran-2-il

Method M

a) By hydration of nitrile B3A (770 mg; 1 mmole) with PtO₂ as a catalyst (130 mg) in absolute ethanol under pressure of 40 atm for 2 days, 750 mg of the raw product was obtained following filtration and evaporation, it was purified on a silica gel column (eluent:chloroform:methanol:ammonia = 6:1:0.1). The quantity of 350 mg of amine M_L1 was obtained.

According to the above procedure, starting from nitrile B4A, amine M_L2 is obtained.

b) The compound **F1A** (282 mg; 0.33 mmole) was dissolved in 3 mL of methanol. Added to the solution was ethylenediamine (0.235 mL; 3.33 mmole) and potassium iodide (581.4 mg; 3.28 mmole). The reaction mixture was stirred at the temperature of 50°C overnight. Methanol was then evaporated under reduced pressure and the residue was dissolved in dichloromethane and washed with water and saturated NaCl solution. After extraction, the organic layer was dried over anhydrous Na₂SO₄ and the solvent was evaporated. The quantity of 240 mg of amine **M_L3** was obtained.

According to the above procedure, starting individually from the compounds **F2A**, **F3A**, **F4A** or **F5A**, amines **M_L4**, **M_L5**, **M_L6**, or **M_L7** are obtained.

According to the above procedure, adding different diamines to the compounds **F2A** and **F5A**, amines **M_L11** and **M_L12** are obtained.

According to the above procedure, by reaction of propylene diamine on the compound **F4A**, amine **M_L8** is obtained.

c) The compound **C1A** (1.27 g; 1.6 mmole) was dissolved in 60 mL of toluene in a flow of argon. After 10 min, 0.672 mL of triethylamine and 0.155 mL of 3-chloropropionylchloride were added. After 15 min, the aliquot of reagents was added again. The reaction mixture was stirred at room temperature for 3 h and water was then added to the reaction mixture. The organic layer was separated, washed with saturated NaHCO₃ solution and dried. After solvent evaporation, the product was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0,1. The quantity of 0.946 g of the compound **M_L9** was obtained.

Similar reaction conditions produce the compound **M_L10**.

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The characteristics of the compounds **M_L1** to **M_L10** are given in Table 4.

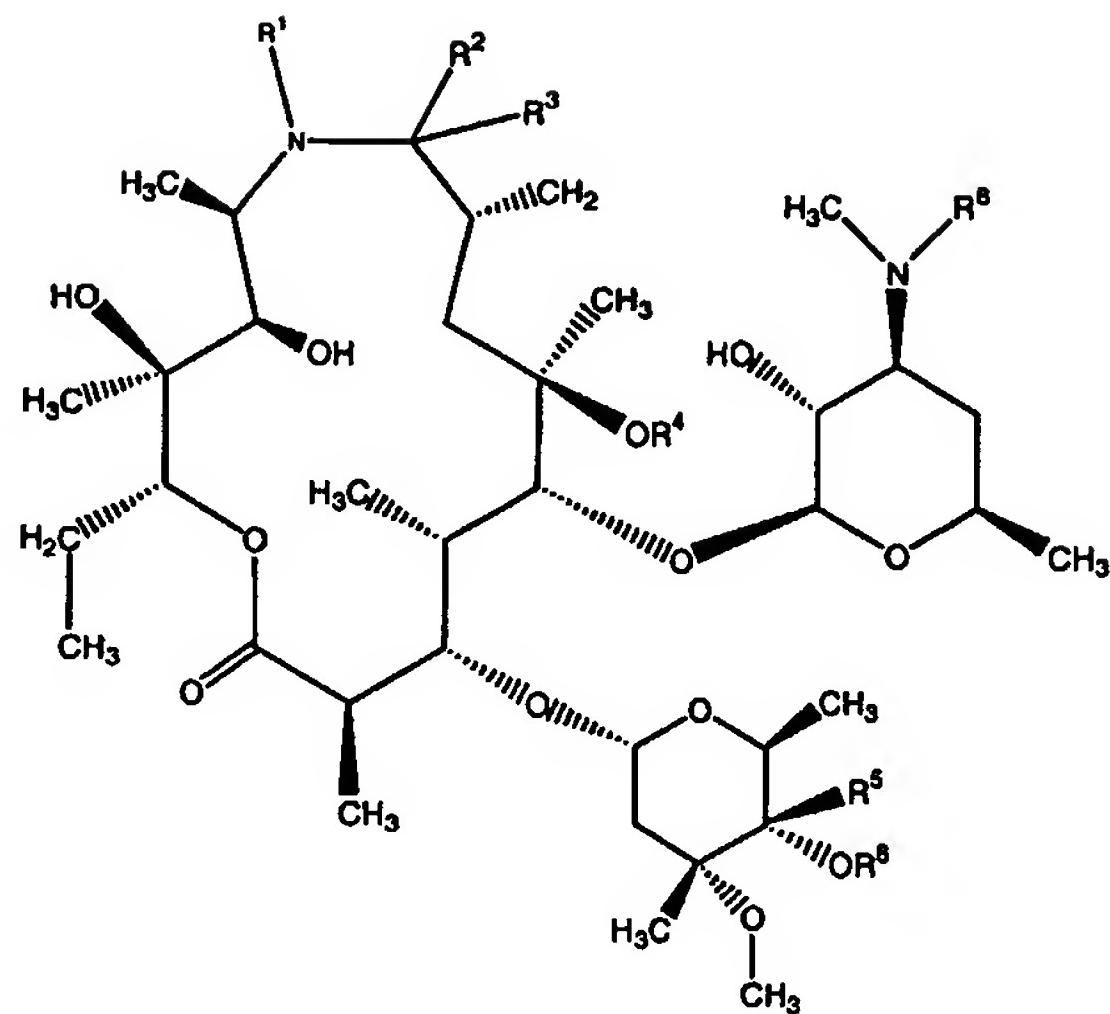
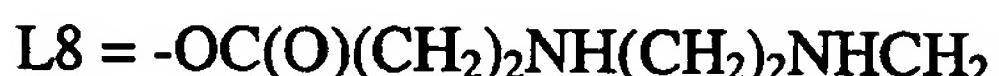
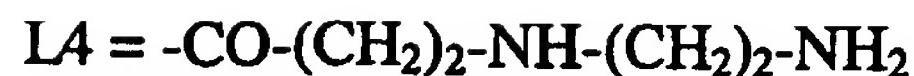


Table 4

Com.	M	R ¹	R ²	R ³	R ⁴	R ⁵	R ⁶	R ⁸	Molecular formula	MH ⁺
M _{L1}	M1	CH ₃	H	H	H	H	H	L1	C ₄₀ H ₇₇ N ₃ O ₁₂	792.6
M _{L2}	M2	H	C=O		CH ₃	H	H	L1	C ₄₀ H ₇₅ N ₃ O ₁₃	806.7
M _{L3}	M1	CH ₃	H	H	H	L2	H	CH ₃	C ₄₁ H ₈₀ N ₄ O ₁₂	821.7
M _{L4}	M2	H	C=O		CH ₃	L2	H	CH ₃	C ₄₁ H ₇₉ N ₄ O ₁₃	835.7
M _{L5}	M3	IPA ^a	H	H	H	L2	H	CH ₃	C ₄₄ H ₈₅ N ₅ O ₁₃	892.7
M _{L6}	M4	CH ₃	H	H	H	L2	H	C ₂ H ₅	C ₄₂ H ₈₂ N ₄ O ₁₂	835.6
M _{L7}	M5	H	C=O		CH ₃	L2	H	C ₂ H ₅	C ₄₂ H ₉₀ N ₄ O ₁₃	849.9
M _{L8}	M4	CH ₃	H	H	H	L3	H	C ₂ H ₅	C ₄₄ H ₈₆ N ₄ O ₁₂	863.8
M _{L9}	M1	CH ₃	H	H	L4	H	H	CH ₃	C ₄₁ H ₇₄ N ₂ O ₁₃	803.6
M _{L10}	M1	CH ₃	H	H	H	H	L4	CH ₃	C ₄₁ H ₇₄ N ₂ O ₁₃	803.9
M _{L11}	M5	H	C=O		CH ₃	L5	H	C ₂ H ₅	C ₄₅ H ₈₆ N ₄ O ₁₃	891.6
M _{L12}	M5	H	C=O		CH ₃	L6	H	C ₂ H ₅	C ₄₈ H ₉₂ N ₄ O ₁₃	933.6

^a IPA = -C(O)-NH-CH(CH₃)₂L1 = -CH₂(CH₂)₂-NH₂L2 = -CH₂-NH-(CH₂)₂-NH₂



L1-L8 corresponds to the link **L**.

Method S

To the solution of the compound **S1** (1.18 g; 3.12 mmole) in 5 mL of DMF were added, *N*-hydroxysuccinimide (NHS) (723 mg ; 6.24 mmol) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (0.598 g; 3.12 mmole). The reaction mixture was stirred at room temperature overnight. Added to the solution were then NH₂CH₂CH₂NH-Boc (0.492 mL; 3.12 mmole) and triethylamine (0.426 mL; 3.062 mmole). The reaction mixture was stirred at room temperature for additional 24 h. The mixture was subsequently transferred to 20 mL of dichloromethane and washed with saturated solution of NaHCO₃ and water. After drying over anhydrous Na₂SO₄, the solvent was evaporated. The quantity of 2.2 g of oily product was obtained; it was dissolved in 30 mL of TFA:CH₂Cl₂ and stirred at room temperature for 1 h. The solvent was subsequently evaporated under reduced pressure. The quantity of 1.24 g of amide **S3** was obtained.

Method Sa

A solution of steroid **S1** (1,0 g, 2,64 mmol) and triethylamine (0,74 ml, 5,29 mmol) in CH₂Cl₂ (60 ml) at 0°C was treated with acryloyl chloride (0,429 ml, 5,29 mmol). After 30 min the reaction mixture was diluted with CH₂Cl₂, washed with aqueous NaHCO₃ and then H₂O, dried and evaporated to give the solid intermediate. This was stirred in acetone (50 ml) with diethylamine (1,38 ml, 13,215 mmol) for

2 hours. Solution was concentrated, diluted with water and washed with EtOAc. The aqueous phase was acidified to pH 2 with 2 N HCl and filtered to provide a solid. 915 mg of compound **S5** was obtained.

Method Sb

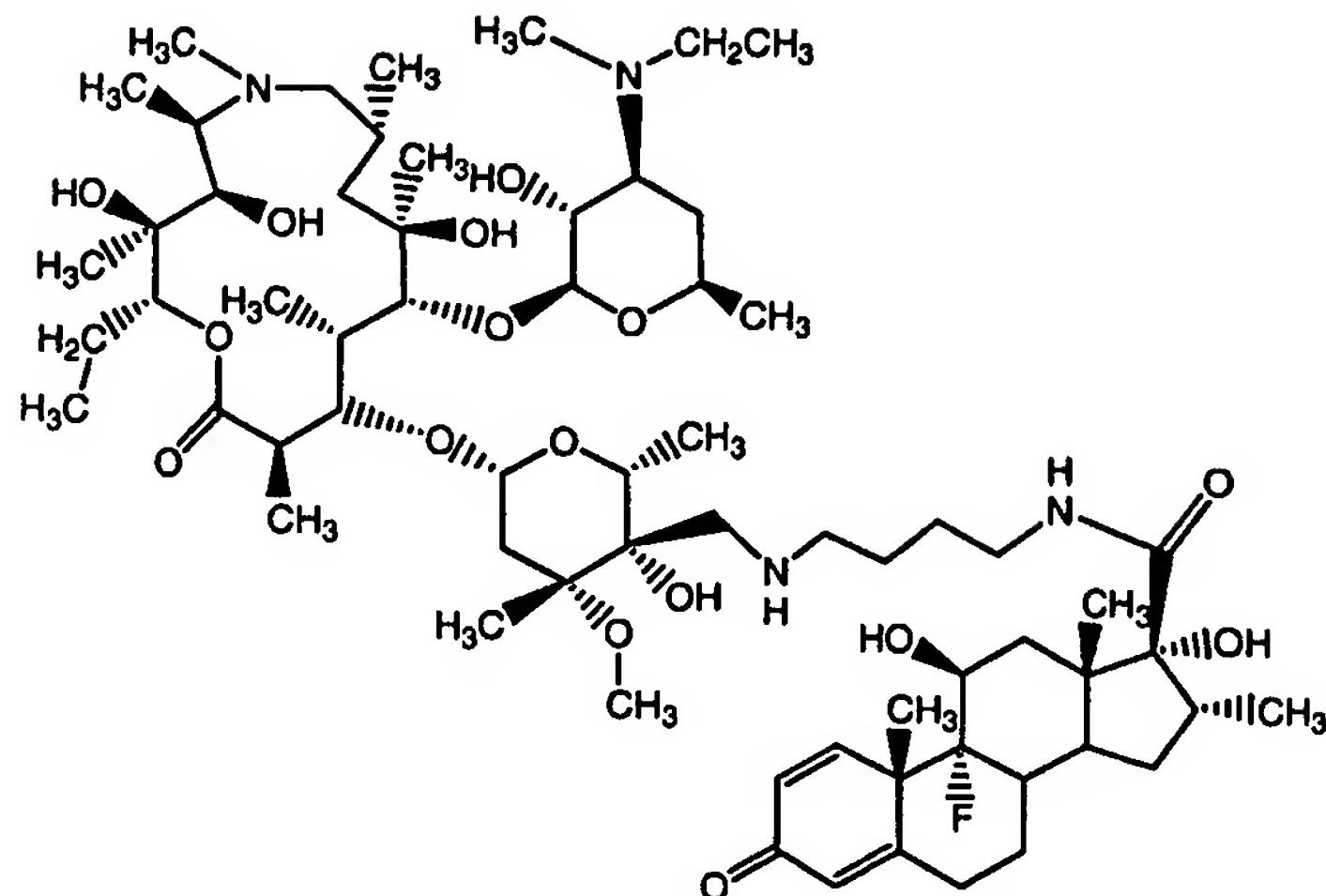
1,8-Diazabicyclo[5.4.0]undec-7-ene (DBU) (1 equiv, 0,23 mmol, 0,035 ml) was added to a 10% solution of acid **S5** (0,23 mmol, 100 mg) in dimethylcarbonate and the resulting mixture was heated to reflux (90°C). After completion, the reaction mixture was cooled to room temperature and diluted with EtOAc and water. The organic layer was dried over Na₂SO₄, evaporated and purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 90:8:1. 80 mg of the compound **S6** was obtained.

Method Sc

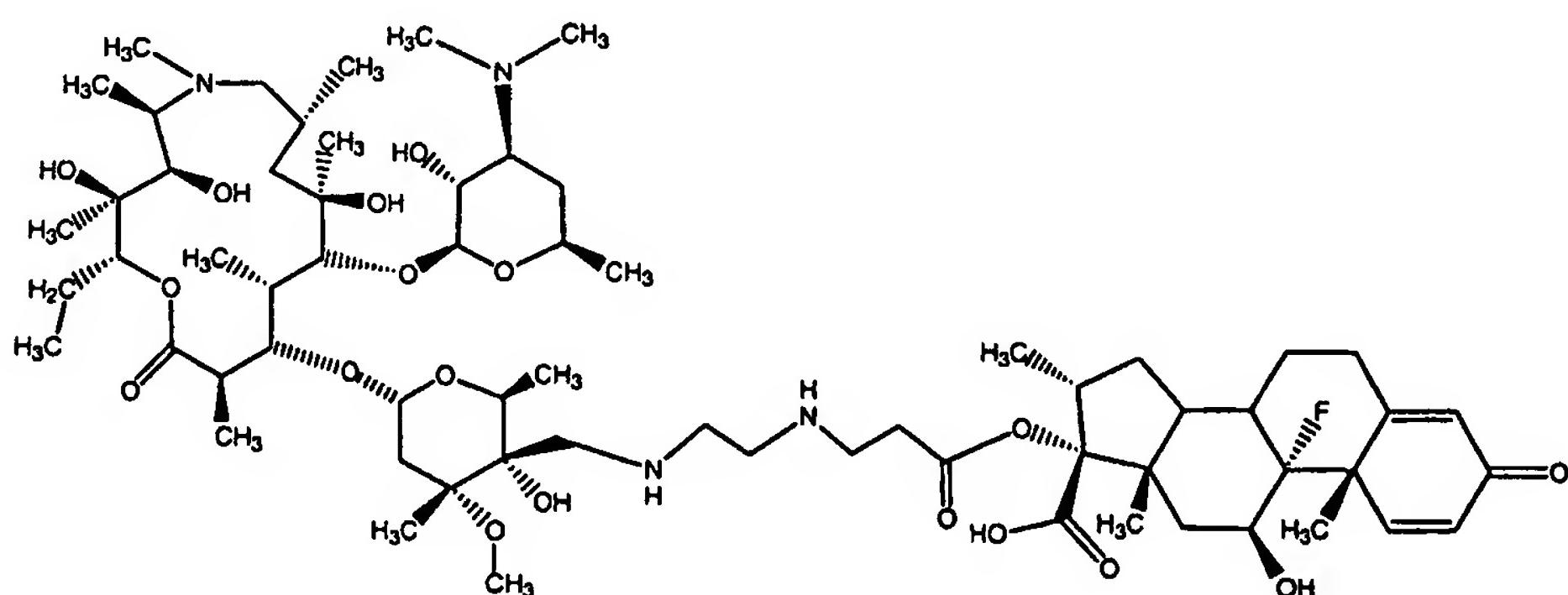
A solution of compound **S5** (184 mg, 0,24 mmol) and N,N-dimethylthiocarbamoyl chloride (60,5 mg, 0,49 mmol) in 2-butanone (8 ml) at room temperature was treated sequentially with triethylamine (0,075 ml, 0,54 mmol), sodium iodide (37 mg, 0,24 mmol), and water (0,018 ml) and stirred for 3 days. Reaction mixture was then treated sequentially with dimethylacetamide (0,88 ml) and water (5,54 ml); cooled to 0°C, stirred for 2 hours and extracted with EtOAc. Organic layer was dried over Na₂SO₄ and evaporated under reduced pressure. Mixture was purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 90:8:1. 20 mg of the compound **S7** was obtained.

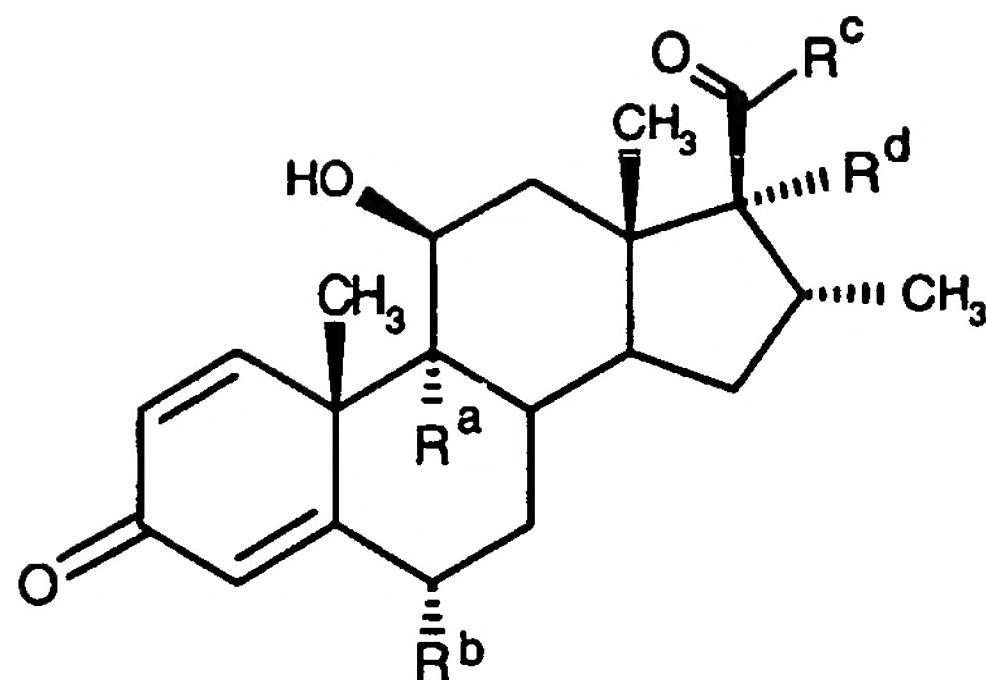
Method Sd

a) In a solution of compound **S5** (250 mg, 0,58 mmol) in methanol (20 ml) 915 mg (1,16 mmol) of the macrolide **A4** was added. Reaction mixture was stirred at 55°C for 24 hours. After evaporation of the solvent, mixture was purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 30:50:2. 540 mg of the compound **S8** was obtained. MS (m/z): 1224,32 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2972, 2939, 2876, 1726, 1686, 1664, 1618, 1605, 1561, 1509, 1459, 1380, 1247, 1168, 1116, 1074, 1056, 1013, 894, 818, 802.



b) In a solution of compound **S5** (250 mg, 0,58 mmol) in methanol (20 ml), 755 mg (1,16 mmol) of the macrolide **M_L3** was added. Reaction mixture was stirred at 55°C for 24 hours. After evaporation of the solvent, mixture was purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 30:50:2. 412 mg of the compound **S9** was obtained. MS (*m/z*): 1253,58 [MH]⁺



**Table 5**

S	R ^a	R ^b	R ^c	R ^d	Molecular formula	MH ⁺
S1	F	H	OH	OH	C ₂₁ H ₂₇ FO ₅	378.9
S2	F	F	OH	OH	C ₂₁ H ₂₆ F ₂ O ₅	397.4
S3	F	H	NH-(CH ₂) ₂ -NH ₂	OH	C ₂₃ H ₃₃ FN ₂ O ₄	421.3
S4	F	H	OH	H	C ₂₁ H ₂₇ FO ₄	363.2
S5	F	H	OH	OC(O)CH ₂ =CH ₂	C ₂₄ H ₂₉ FO ₆	433.2
S6	F	H	CH ₃	OC(O)CH ₂ =CH ₂	C ₂₅ H ₃₁ FO ₆	446.9
S7	F	H	SC(O)N(CH ₃) ₂	OC(O)CH ₂ =CH ₂	C ₂₇ H ₃₄ FNO ₆ S	520.2

Example 23

Compound 23: I; M = M₁, L = L₁, S = S₁; (R¹ = R⁷ = CH₃, R² = R³ = R⁴ = R⁵ = R⁶ = H, X¹ = CH₂, m = 2, Q = NH, n = 0, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound **S1** (84 mg; 0.22 mmole) was dissolved in 5 mL of dry dichlormethane. To the solution, 0.29 mL of triethylamine and 61 mg of hydroxybenzotriazole were added and subsequently the macrolide **M_{L1}** (175 mg; 0.22 mmole) and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (179 mg) were added. The reaction mixture was stirred at room temperature overnight. The solvent was evaporated under reduced pressure and the obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH = 6:1:0.1. 16 mg of compound **23** was obtained; MS (*m/z*): 1152.5 [MH]⁺.

Example 24

Compound 24: I; M = M₂, L = L₁, S = S₁; (R¹ = R⁵ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, X¹ = CH₂, m = 2, Q = NH, n = 0, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound **S1** (117.5 mg; 0.31 mmole) was dissolved in 5 mL of dry DMF. To the solution, 58.6 µl of diisopropylethylamine and 42 mg of 1-hydroxybenzotriazole were added and then the macrolide **M_{L2}** (250 mg; 0.31 mmole) and 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (60 mg) were added. The reaction mixture was stirred at the temperature of 100°C for 5 hours. The mixture was extracted twice with 40 mL of ethyl-acetate and 40 mL of water each time. The organic layers were washed twice with 30 mL of water each time and dried over anhydrous sodium-sulphate. The solvent was evaporated under reduced pressure and the obtained mixture was purified on a silica gel column, eluent CHCl₃:MeOH:NH₄OH = 6:1:0.1, with 65 mg of compound **24** being obtained; MS (*m/z*): 1166.7 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2972, 2938, 2881, 1774, 1736, 1702, 1664, 1560, 1528, 1459, 1379, 1298, 1244, 1170, 1126, 1108, 1075, 1055, 1011, 959, 894.

Example 25

Compound 25: I; M = M1, L = L2, S = S2; (R¹ = R⁷ = R⁸ = CH₃, R² = R³ = R⁴ = R⁶ = H, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = R^b = F, R^d = OH)

The compound **S2** (116 mg; 0.29 mmole) was dissolved in 10 mL of dichloromethane. Added were 0.320 mL of triethylamine and 78 mg of 1-hydroxybenzotriazole and then the compound **M_L3** (240 mg; 0.29 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (200 mg). The reaction mixture was stirred at room temperature in an inert atmosphere for 24 h. The solvent was then evaporated and the compound purified on a silica gel column in the solvent system CHCl₃: MeOH:NH₄OH = 6:1:0.1. 214 mg of compound **25** was isolated; MS (*m/z*): 1200.0 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2973, 2939, 2876, 1711, 1670, 1632, 1561, 1523, 1458, 1380, 1317, 1259, 1177, 1130, 1110, 1073, 1034, 995, 939, 898, 795, 757, 710, 642.

Example 26

Compound 26: I; M = M1, L = L2, S = S1; (R¹ = R⁷ = R⁸ = CH₃, R² = R³ = R⁴ = R⁶ = H, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = OH)

To the suspension of the compound **S1** (111 mg; 0.29 mmole) in dry CH₂Cl₂ (5 mL) cooled to 0°C under argon were added, 0.380 mL of triethylamine and 80 mg of 1-hydroxybenzotriazole and then the compound **M_L3** (240 mg; 0.29 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (235 mg). The reaction mixture was stirred at room temperature in a flow of argon for 24 hours and then evaporated to a smaller volume under reduced pressure and purified on a silica gel column (eluent: CHCl₃:CH₃OH:NH₄OH = 6:1:0.1). 88 mg of compound **26** was obtained; MS (*m/z*): 1182 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2971, 2939, 2874, 1719, 1665, 1625, 1560, 1518, 1458, 1378, 1275, 1175, 1110, 1036, 1013, 996, 894, 796, 757, 670.

Example 27

Compound 27: I; M = M2, L = L2, S = S2; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = R⁸ = CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = R^b = F, R^d = OH)

The compound **S2** (47 mg; 0.12 mmole) was dissolved in dry CH₂Cl₂ (5 mL) in an inert atmosphere and 0.128 mL of triethylamine and 31 mg of 1-hydroxybenzotriazole were added, subsequently, the compound **M_L4** (141 mg; 0.12 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (169 mg) were added. The reaction mixture was stirred at room temperature in a flow of argon for 24 hours and then evaporated to a smaller volume under reduced pressure and purified on a silica gel column (eluent: CHCl₃:CH₃OH:NH₄OH = 6:1:0.1). 82 mg of compound **27** was obtained; MS (*m/z*): 1213 [MH]⁺.

Example 28

Compound 28: I; M = M3, L = L2, S = S2; R¹ = CONHCH(CH₃)₂, R² = R³ = R⁴ = R⁶ = H, R⁷ = R⁸ = CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = R^b = F, R^d = OH)

The compound **S2** (95 mg; 0.24 mmole) was dissolved in 10 mL of dichloromethane, 0.262 mL of triethylamine and 65 mg of 1-hydroxybenzotriazole were added and then the compound **M_L5** (257 mg; 0.24 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (165 mg) were added. The reaction mixture was stirred at room temperature in an inert atmosphere for 24 h. The solvent was then evaporated and the compound purified on a silica gel column in the solvent system CHCl₃: MeOH:NH₄OH = 6:1:0.1. 144 mg of compound **28** was isolated; MS (*m/z*): 1270 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2974, 2940, 2877, 1774, 1735, 1719, 1670, 1630, 1561, 1523, 1459, 1381, 1317, 1264, 1168, 1110, 1072, 1032, 994, 899, 820, 757, 709.

Example 29

Compound 29: I; M = M4, L = L2, S = S1; (R¹ = R⁷ = CH₃, R² = R³ = R⁴ = R⁶ = H, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound **S1** (282 mg; 0.71 mmole) was dissolved in dry CH₂Cl₂ (15 mL) in an inert atmosphere, 0.776 mL of triethylamine and 192 mg of 1-hydroxybenzotriazole were added and then the compound **M_L6** (593 mg; 2.85 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (489 mg) were added. The reaction mixture was stirred in a flow of argon at room temperature for 24 hours and then evaporated to a smaller volume under reduced pressure and purified on a silica gel column (eluent: CHCl₃:CH₃OH:NH₄OH = 6:1:0.1). 84 mg of compound **29** was obtained; MS (*m/z*): 1195.9 [MH]⁺. IR(cm⁻¹)/KBr: 3424, 2972, 2939, 2874, 1725, 1665, 1625, 1560, 1522, 1459, 1379, 1259, 1176, 1109, 1054, 1036, 1012, 996, 894, 796, 757.

Example 30:

Compound 30: I; M = M5, L = L2, S = S1; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = OH)

In 15 mL of dry dichloromethane, the compound **S1** (89 mg; 0.23 mmole) was dissolved. Added were 0.257 mL of triethylamine and 64 mg of 1-hydroxybenzotriazole and then the compound **M_L7** (200 mg; 0.23 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (162 mg). The reaction mixture was stirred in an inert atmosphere at room temperature for 24 h. The solvent was then evaporated under reduced pressure and the compound purified in a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 153 mg of compound **30** was obtained; MS (*m/z*): 1209.7 [MH]⁺. IR(cm⁻¹)/KBr: 3423, 2974, 2939, 2881, 1774, 1708, 1664, 1560, 1523, 1459, 1379, 1299, 1275, 1221, 1175, 1109, 1053, 1010, 893, 813, 757.

Example 31

Compound 31: I; M = M5, L = L2, S = S4; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = H)

The compound S4 (85 mg; 0.24 mmole) was dissolved in 10 mL of dry dichloromethane. Added were 0.257 mL of triethylamine and 64 mg of 1-hydroxybenzotriazole and then the compound M_L7 (200 mg; 0.24 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (162 mg). The reaction mixture was stirred in an inert atmosphere at room temperature for 24 h. The solvent was then evaporated under reduced pressure and the compound was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 106 mg of compound 31 was obtained; MS (*m/z*): 1193.6 [MH]⁺. IR(cm⁻¹)/KBr: 3448, 2973, 2940, 2876, 1708, 1664, 1560, 1528, 1459, 1379, 1297, 1272, 1225, 1176, 1110, 1053, 1010, 929, 889, 810, 756.

Example 32

Compound 32: I; M = M4, L = L3, S = S1; (R¹ = R⁷ = CH₃, R² = R³ = R⁴ = R⁶ = H, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 4, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound S1 (74 mg; 0.19 mmole) was dissolved in dry CH₂Cl₂ (10 mL) in an inert atmosphere and 0.176 mL of triethylamine and 53 mg of 1-hydroxybenzotriazole were added, followed by the addition of the compound M_L8 (169 mg; 0.19 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (135 mg). The reaction mixture was stirred in a flow of argon at room temperature for 24 hours and then evaporated to a smaller volume under reduced pressure and purified on a silica gel column (eluent: CHCl₃:CH₃OH:NH₄OH = 6:1:0.1). 67 mg of compound 32 was obtained; MS (*m/z*): 1223.6 [MH]⁺. IR(cm⁻¹)/KBr: 3425, 2979, 2937, 2873, 2370, 1734, 1665, 1627, 1562, 1525, 1459, 1379, 1259, 1175, 1109, 1054, 1036, 1012, 995, 893, 796, 756.

Example 33

Compound 33: I; M = M1, L = L5, S = S1; (R¹ = R⁷ = R⁸ = CH₃, R² = R³ = R⁴ = R⁵ = H, X¹ = CO, m = 2, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound **S3** (109 mg; 0.26 mmole) was dissolved in 20 mL of methanol and then the compound **M_L10** (250 mg; 0.31 mmole) was added. The reaction mixture was stirred at 75 °C for 24 h.. The solvent was then evaporated under reduced pressure and the compound was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 40 mg of compound **33** was obtained; MS (*m/z*): 1223.9 [MH]⁺. IR(cm⁻¹)/KBr: 3424, 2972, 2939, 2873, 1734, 1665, 2639, 1562, 1525, 1459, 1379, 1260, 1173, 1108, 1074, 1036, 1015, 960, 894, 835, 797, 755, 642.

Example 34

Compound 34: I; M = M1, L = L5, S = S1; (R¹ = R⁷ = R⁸ = CH₃, R² = R³ = R⁵ = R⁶ = H, X¹ = CO, m = 2, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d = OH)

The compound **M_L9** (267 mg; 0.32 mmole) was dissolved in 30 mL of methanol. To the solution, the compound **S3** (399 mg; 0.95 mmole) was added. The reaction mixture was stirred at 75 °C overnight. The solvent was then evaporated and the obtained mixture was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 30 mg of compound **34** was obtained; MS (*m/z*): 1223.7 [MH]⁺. IR(cm⁻¹)/KBr: 345, 2972, 2941, 2873, 1722, 1665, 1627, 1525, 1459, 1379, 1296, 1243, 1184, 1110, 1053, 1012, 894, 831, 756.

Example 35

Compound 35: I; M = M5, L = L6, S = S4; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 5, X² = NH, R^a = F, R^b = H, R^d = H)

The compound **S4** (63 mg; 0.17 mmole) was dissolved in 10 mL of dry dichloromethane. Added were 0.188 mL of triethylamine and 47 mg of 1-

hydroxybenzotriazole and then the compound **M_L11** (154 mg; 0.17 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (119 mg). The reaction mixture was stirred in an inert atmosphere at room temperature for 24 h. The solvent was then evaporated under reduced pressure and the compound was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 250 mg of compound **35** was obtained; MS (*m/z*): 1236,5 [MH]⁺. IR(cm⁻¹)/KBr: 3448, 2972, 2938, 2870, 1736, 1708, 1664, 1560, 1535, 1459, 1379, 1329, 1297, 1272, 1229, 1176, 1110, 1053, 1010, 931, 889, 809.

Example 36

Compound 36: I; M =M5, L = L7, S =S4; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 8, X² = NH, R^a = F, R^b = H, R^d=H)

The compound **S4** (73 mg; 0.20 mmole) was dissolved in 10 mL of dry dichloromethane. Added were 0.219 mL of triethylamine and 54 mg of 1-hydroxybenzotriazole and then the compound **M_L12** (188 mg; 0.20 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (139 mg). The reaction mixture was stirred in an inert atmosphere at room temperature for 24 h. The solvent was then evaporated under reduced pressure and the compound was purified on a silica gel column in the solvent system CHCl₃:MeOH:NH₄OH = 6:1:0.1. 54 mg of compound **36** was obtained; MS (*m/z*): 1277,6 [MH]⁺.

Example 37

Compound 37: I; M =M2, L = L6, S =S4; (R¹ = R⁶ = H, R² together with R³ = CO, R⁴ = R⁷ = CH₃, R⁸ = CH₂CH₃, X¹ = CH₂, m = 0, Q = NH, n = 2, X² = NH, R^a = F, R^b = H, R^d=H)

The compound **S4** (34 mg; 0.093 mmole) was dissolved in 10 mL of dry dichloromethane. Added were 0.100 mL of triethylamine and 25 mg of 1-hydroxybenzotriazole and then the compound **M_L4** (29 mg; 0.093 mmole) and 1-(3-dimethylaminopropyl)-3-ethyl-carbodiimide hydrochloride (64 mg). The reaction

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mixture was stirred in an inert atmosphere at room temperature for 24 h. The solvent was then evaporated under reduced pressure and the compound was purified on a silica gel column in the solvent system $\text{CHCl}_3:\text{MeOH}:\text{NH}_4\text{OH} = 6:1:0.1$. 10 mg of compound 37 was obtained; MS (m/z): 1179,7 $[\text{MH}]^+$.

Example 38

Compound 38: I; M = M1, L = L7, S = S6; (R² = R³ = R⁴ = R⁵ = R⁶ = H, R⁸ = CH₃, X₁ = -CH₂-, m = 2, n = 2 Q = -NH-, X² = OC(O), R^a = F, R^b = H, R^c = OCH₃)

a) 1, 8-Diazabicyclo[5.4.0]undec-7-ene (DBU) (1 equiv, 0,082 mmol, 0,012 ml) was added to a 10% solution of acid S8 (100 mg, 0,082 mmol) in dimethylcarbonate (1 ml) and the resulting mixture was heated to reflux (90°C). After completion, the reaction mixture was cooled to room temperature and diluted with EtOAc and water. The organic layer was dried over Na₂SO₄, evaporated and purified on a silica gel column in the solvent system $\text{CH}_2\text{Cl}_2:\text{MeOH}:\text{NH}_4\text{OH} = 90:8:1$. 32 mg of the compound 38 was obtained. MS (m/z): 1238,55 $[\text{MH}]^+$. IR(cm^{-1})/KBr: 3449, 2972, 2939, 2877, 1736, 1666, 1625, 1561, 1542, 1509, 1459, 1377, 1291, 1266, 1241, 1176, 1115, 1052, 1014, 978, 957, 893, 833, 812, 755, 705, 639.

b) In a solution of compound S6 (17,3 mg, 0,039 mmol) in methanol (3 ml) 62 mg (0,094 mmol) of the macrolide A4 was added. Reaction mixture was stirred at 55°C for 24 hours. After evaporation of the solvent, mixture was purified on a silica gel column in the solvent system $\text{CH}_2\text{Cl}_2:\text{MeOH}:\text{NH}_4\text{OH} = 90:8:1$. 35 mg of the compound 38 was obtained. MS (m/z): 1238,55 $[\text{MH}]^+$

Example 39

Compound 39: I; M = M1, L = L8, S = S6; (R¹ = R⁷ = R⁸ = CH₃, R² = R³ = R⁴ = R⁶ = H, X¹ = CH₂NH, m = 2, Q = NH, n = 2, X² = OC(O), R^a = F, R^b = H, R^c = OCH₃)

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In a solution of compound **S6** (100 mg, 0,22 mmol) in methanol (10 ml), 212 mg (0,26 mmol) of the macrolide **M_L3** was added. Reaction mixture was stirred at 55°C for 24 hours. After evaporation of the solvent, mixture was purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 90:8:1. 20 mg of the compound **39** was obtained. MS (*m/z*): 1267,55 [MH]⁺

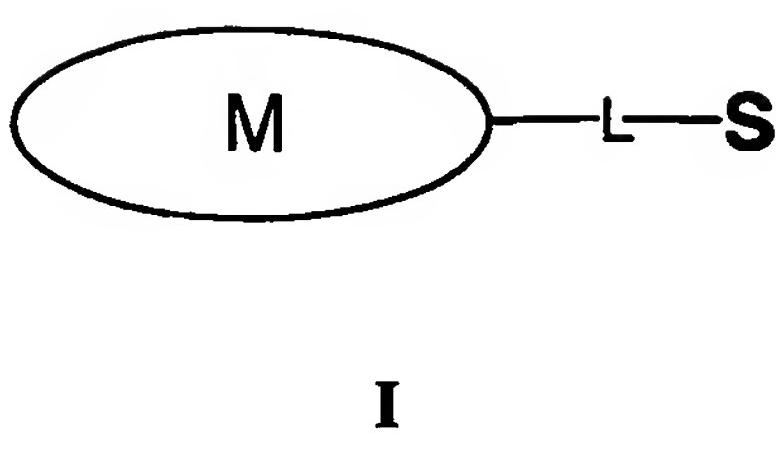
Example 40

Compound 40: I; M =M1, L = L7, S =S7; (R² = R³ = R⁴ = R⁵ = R⁶ =H, R⁸ = CH₃, X1 = -CH₂-, m = 2, n = 2 Q = -NH-, X² = OC(O), R^a = F, R^b = H, R^c = SC(O)N(CH₃)₂)

A solution of compound **S8** (150 mg, 0,12 mmol) in 2-butanone (4 ml) at room temperature was treated sequentially with triethylamine (37,5 ml, 0,27 mmol), sodium iodide (18,4 m, 0,12 mmol), and water (0,015 ml) and stirred for 3 days. Reaction mixture was then treated sequentially with dimethylacetamide (0,44 ml) and water (2,77 ml); cooled to 0°C, stirred for 2 hours and extracted with EtOAc. Organic layer was dried over Na₂SO₄ and evaporated under reduced pressure. Mixture was purified on a silica gel column in the solvent system CH₂Cl₂:MeOH:NH₄OH = 90:8:1. 56,3 mg of the compound **40** was obtained. MS (*m/z*): 1311,49 [MH]⁺

CLAIMS

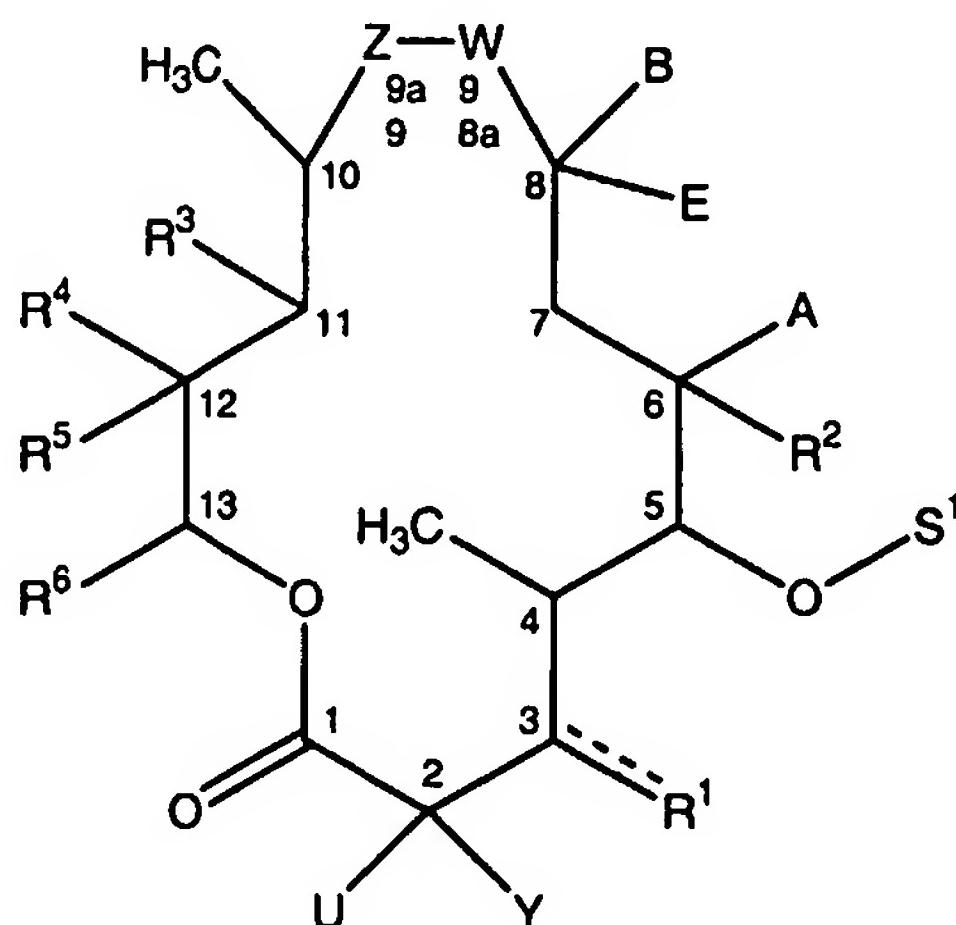
1. A compound of the formula:



wherein

- (i) **M** is a macrolide subunit selected from the group consisting of multi-member lactonic ring molecules, wherein "member" refers to the number of carbon atoms or heteroatoms in the lactonic ring and "multi" signifies a whole number greater than about 10, and up to about 18 said molecules having the property of accumulating within mammalian, including human, immune system cells that mediate inflammatory immune responses;
- (ii) **S** is a steroidal anti-inflammatory subunit; and
- (iii) **L** is a linker molecule to which each of **M** and **S** are covalently linked.

2. A compound as claimed in claim 1 wherein **M** represents a group of Formula II:



wherein

- (i) Z and W independently are $>\text{C}=\text{O}$, $>\text{CH}_2$, $>\text{CH}-\text{NR}_t\text{R}_s$, $>\text{N}-\text{R}_N$ or $>\text{C}=\text{N}-\text{R}_M$, wherein

R_t and R_s independently are hydrogen or alkyl;

R_M is hydroxy, alkoxy, substituted alkoxy or OR^P ;

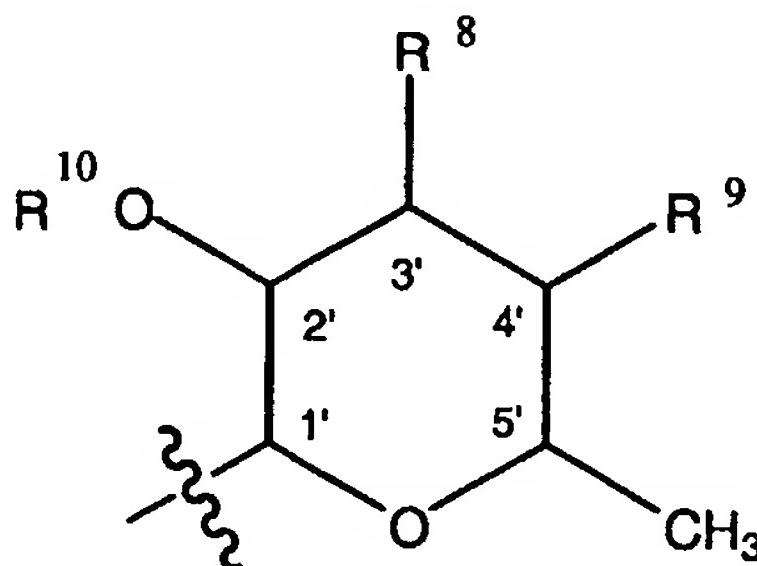
R_N is hydrogen, R^P , alkyl, alkenyl, alkynyl, alkoxy, alkoxyalkyl, or $-\text{C}(\text{X})-\text{NR}_t\text{R}_s$; wherein X is =O or =S;

provided that Z and W cannot both simultaneously be, $>\text{C}=\text{O}$, $>\text{CH}_2$, $>\text{CH}-\text{NR}_t\text{R}_s$, $>\text{N}-\text{R}_N$, $>\text{C}=\text{N}-\text{R}_M$ or a bond;

- (ii) U and Y independently are hydrogen, halogen, alkyl, or hydroxyalkyl;

- (iii) R^1 is hydroxy, OR^P , $-\text{O}-\text{S}^2$ group or an =O;

- (iv) S^1 is a sugar moiety of Formula III:



III

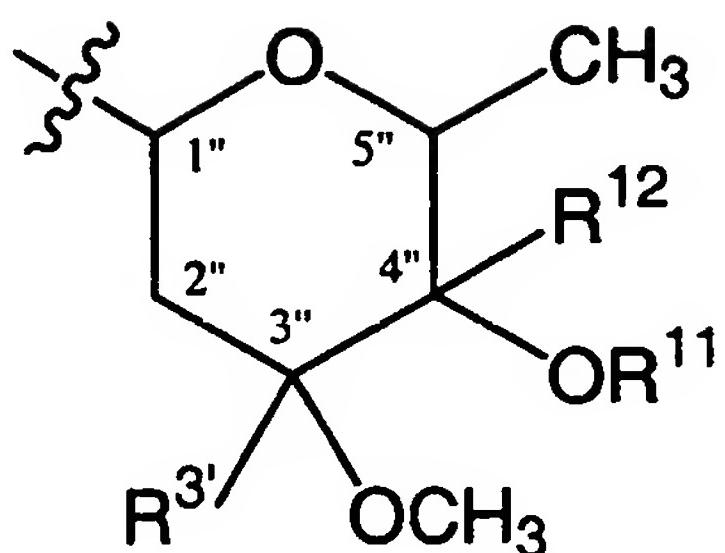
wherein

R^8 and R^9 are both hydrogen or together form a bond, or R^9 is hydrogen and R^8 is $-\text{N}(\text{CH}_3)\text{R}^y$, wherein

R^y is R^P , R^z or $-\text{C}(\text{O})\text{R}^z$, wherein R^z is hydrogen or alkyl or alkenyl or alkynyl or cycloalkyl or aryl or heteroaryl or alkyl substituted with $\text{C}_2\text{-C}_7$ -alkyl, $\text{C}_2\text{-C}_7$ -alkenyl, $\text{C}_2\text{-C}_7$ -alkynyl, aryl or heteroaryl;

R^{10} is hydrogen or R^P ;

- (v) S^2 sugar moiety of Formula IV:



IV

wherein

$R^{3'}$ is hydrogen or methyl;

R^{11} is hydrogen, R^p , or $O-R^{11}$ is a group that with R^{12} and with C/4" carbon atom forms a $>C=O$ or epoxy group;

R^{12} is hydrogen or a group that with $O-R^{11}$ group and with C/4" carbon atom forms a $>C=O$ or epoxy group;

(vi) R^2 is hydrogen, hydroxy, OR^p or alkoxy ;

(vii) A is hydrogen or methyl;

(viii) B is methyl or epoxy;

(ix) E is hydrogen or halogen;

(x) R^3 is hydroxy, OR^p , alkoxy or R^3 is a group that with R^5 and with C/11 and C/12 carbon atoms forms a cyclic carbonate or carbamate, or if W or Z is $>N-R_N$ R^3 is a group that with W or Z forms a cyclic carbamate;

(xi) R^4 is C_1-C_4 alkyl;

(xii) R^5 is hydrogen, hydroxy, OR^p , C_1-C_4 alkoxy, or a group that with R^3 and with C/11 and C/12 carbon atoms forms a cyclic carbonate or carbamate;

(xiii) R^6 is hydrogen or C_1-C_4 -alkyl;

wherein M has a linkage site through which it is linked to S via linking group L; provided that the linkage site being at one or more of the following:

a) any reactive hydroxy, nitrogen, or epoxy group located on S^1 , S^2 , or an aglycone oxygen if S^1 and/or S^2 is cleaved off;

b) a reactive $>N-R_N$ or $-NR_tR_s$ or oxo group located on Z or W;

c) a reactive hydroxy group located at any one of R^1 , R^2 , R^3 , and R^5 ;

d) any other group that can be first derivatized to a hydroxy or $-NR_tR_s$ group and

R^P is hydroxyl or amino protective group.

3. A compound as claimed in claim 1 wherein L represents a group of Formula VA or of Formula VB:

VA	$X^1-(CH_2)_m-X^2$
VB	$X^1-(CH_2)_m-Q-(CH_2)_n-X^2$

wherein

X^1 is selected from: $-CH_2-$, $-CH_2NH-$, $-C(O)-$, $-OC(O)-$, $=N-O-$, $-OC(O)NH-$; or $-C(O)NH$;

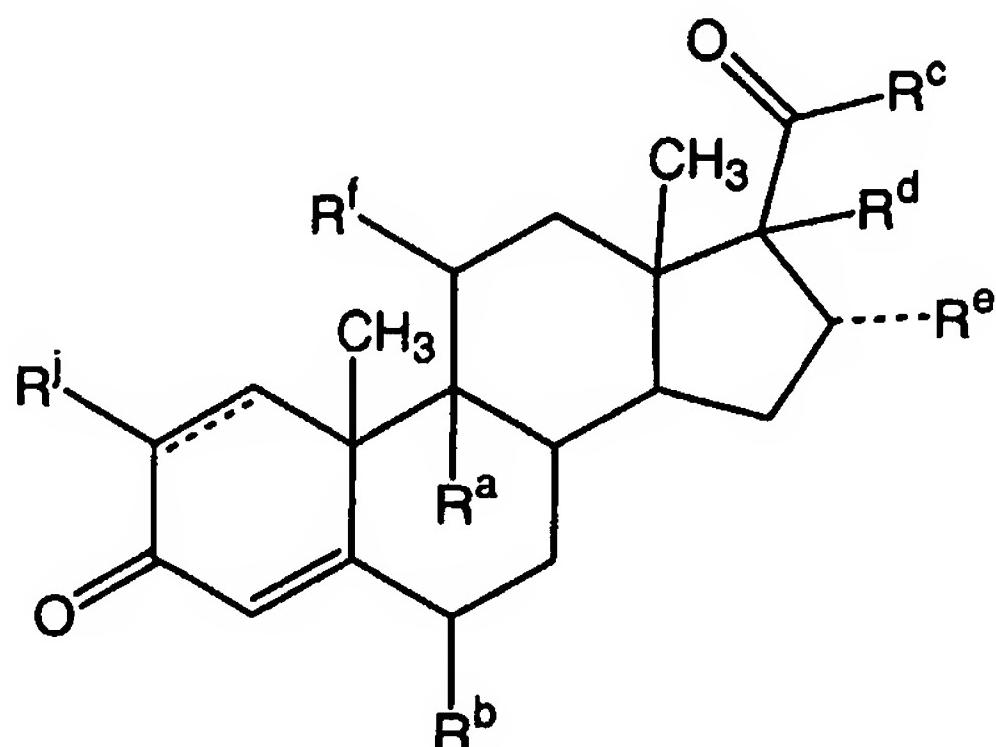
X^2 is $-NH-$, $-NHC(O)-$ or $-CH_2-$;

Q is $-NH-$ or $-CH_2-$, wherein

each $-CH_2-$ or $-NH-$ group may be optionally substituted by C_1-C_7 -alkyl, C_2-C_7 -alkenyl, C_2-C_7 -alkynyl, $C(O)R^x$, $C(O)OR^x$, $C(O)NHR^x$, wherein R^x may be C_1-C_7 -alkyl, aryl or heteroaryl;

the symbols m and n independently are a whole number from 0 to 8, with the proviso that if Q is NH, n cannot be 0.

4. A compound as claimed in claim 1 wherein S represents a group of Formula X:



X

wherein

R^a and R^b independently represents, hydrogen or halogen;

R^c is hydroxy, alkoxy (preferably metoxy), alkyl, thiocarbamoyl, carbamoyl or a valence-bond;

R^d and R^e independently represents: hydrogen, hydroxy, methyl or C₁-C₄-alkoxy or each are a group that forms a 1,3-dioxolane ring with the other or a valence bond;

R^f is hydrogen, hydroxy, chloro, or forming a keto group with the carbon atom it is attached to;

R^j is hydrogen or halogen.

5. A compound as claimed in claim 2 wherein

Z is >NR_N, wherein R_N is hydrogen or a methyl group;

W is >CH₂;

B is methyl;

E is hydrogen;

R² is hydroxy;

A is methyl;

S¹ group represents a group of Formula III wherein

R⁸ is selected from: hydrogen, amino, N-methylamino, N,N-dimethylamino, N-methyl-N-(C₂-C₄)-alkylamino, N-methyl-N-methylcarbonylamino, N-methyl-N-benzylamino, N-methyl-N-cyclohexylamino;

R⁹ and R¹⁰ are hydrogen;

R¹ is O-S² wherein S² represents a group of Formula IV wherein R¹¹ and R¹² are hydrogen and R¹³ is methyl;

U is hydrogen;

Y is methyl;

R⁴ is methyl;

R⁶ is ethyl;

R⁵ is hydroxy or a group that with R³ and with C/11 and C/12 carbon atoms forms a cyclic carbonate bridge;

R^3 is hydroxy or a group that with R^5 and with C/11 and C/12 carbon atoms forms a cyclic carbonate bridge;

provided that the linkage is through the nitrogen of Z at N/9a position or through the oxygen of R^3 at C/11 position.

6. A compound as claimed in claim 2 wherein

Z is selected from >N-H, >N-CH₃, >N-C(O)NHR^x, wherein R^x is isopropyl;

W is >C=O or >CH₂ provided that when Z is >N-CH₃ W cannot be >C=O;

B is methyl;

E is hydrogen;

A is methyl;

R^2 is hydroxy or methoxy;

S^1 group represents a group of Formula III wherein

R^8 is selected from: amino, C₁-C₆-alkylamino, C₁-C₆-dialkylamino;

R^9 and R^{10} are hydrogen;

R^1 is O-S² wherein S² represents a group of Formula IV wherein R^{11} is hydrogen or

O-R¹¹ is a group that with R^{12} and with C/4"carbon atom forms a >C=O or

epoxy group; R^{12} is hydrogen or a group that with O-R¹¹ group and with

C/4"carbon atom forms a >C=O or epoxy group; R^{13} is methyl;

U is hydrogen;

Y is methyl;

R^3 is hydroxy;

R^4 is methyl;

R^5 is hydroxy or methoxy;

R^6 is ethyl;

provided that the linkage is through the nitrogen of R^8 at C/3', through the oxygen of R^2 at C/6 or through the carbon of R^{12} or through the oxygen of R^{11} both at C/4".

7. A compound as claimed in claim 4 wherein

R^a and R^b independently represents, hydrogen or halogen;

R^d is hydrogen or hydroxy;

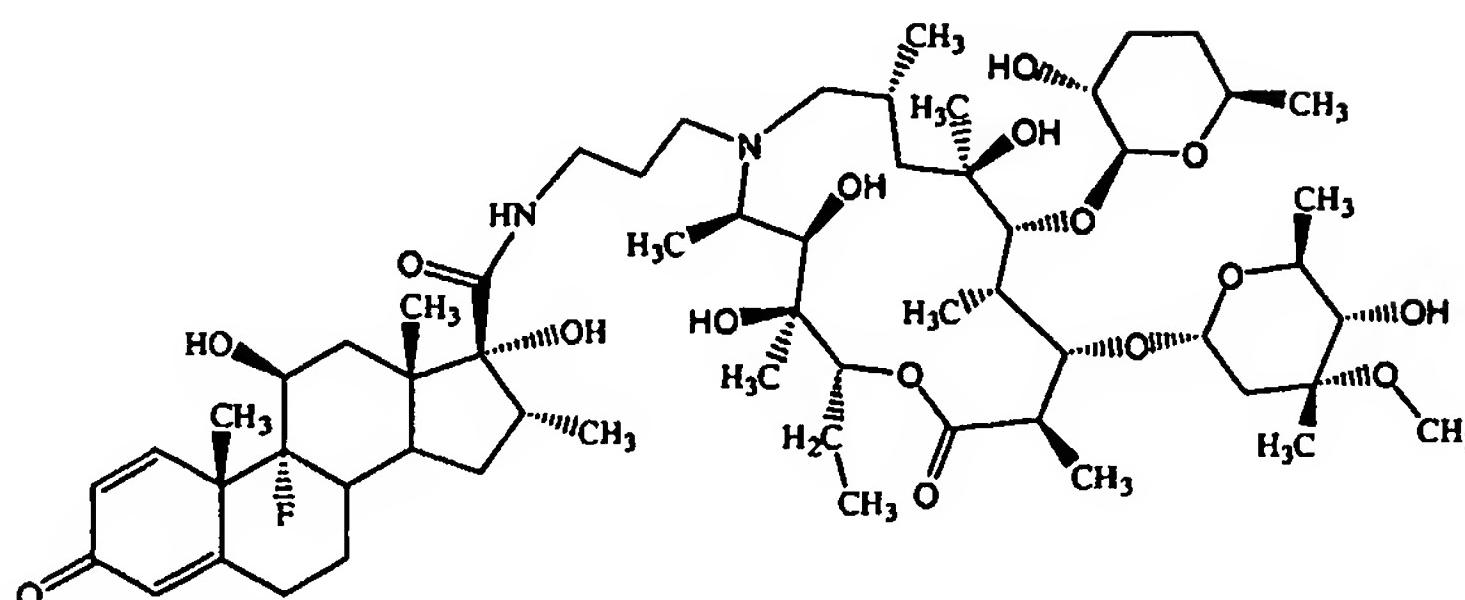
R^e is methyl;

R^f is hydroxy;

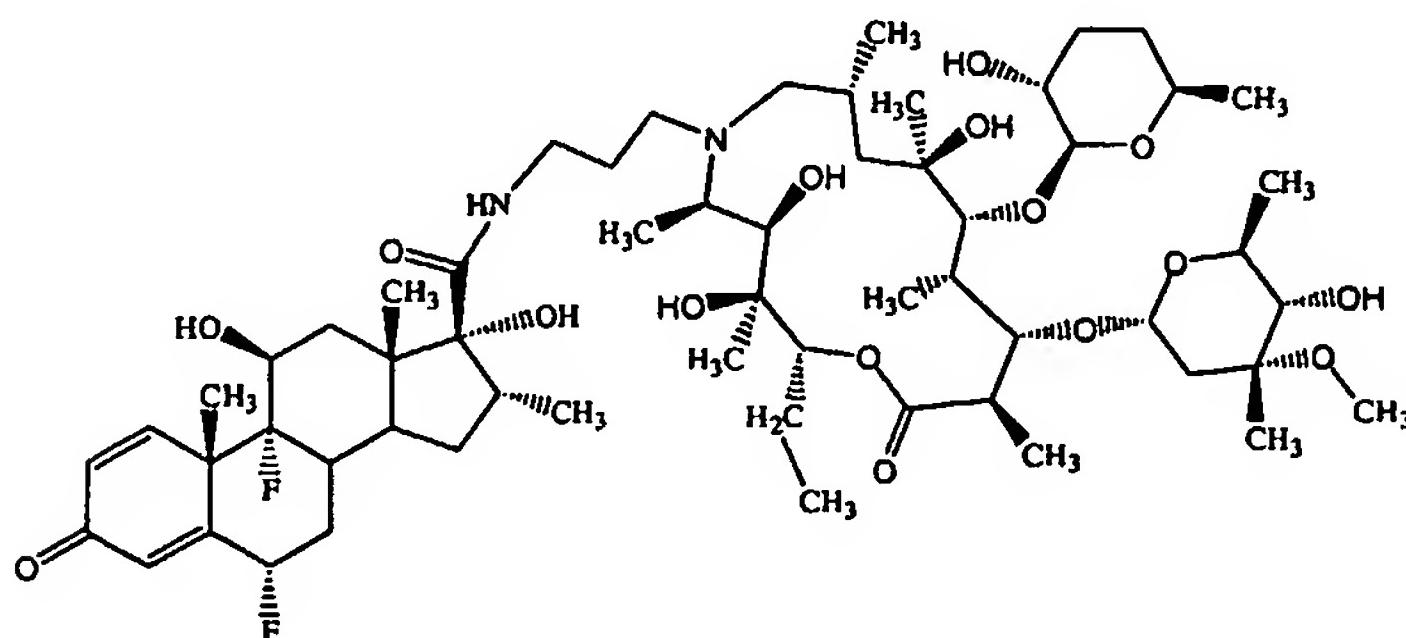
Rj is hydrogen

provided that the linkage is through the valence bond R^k.

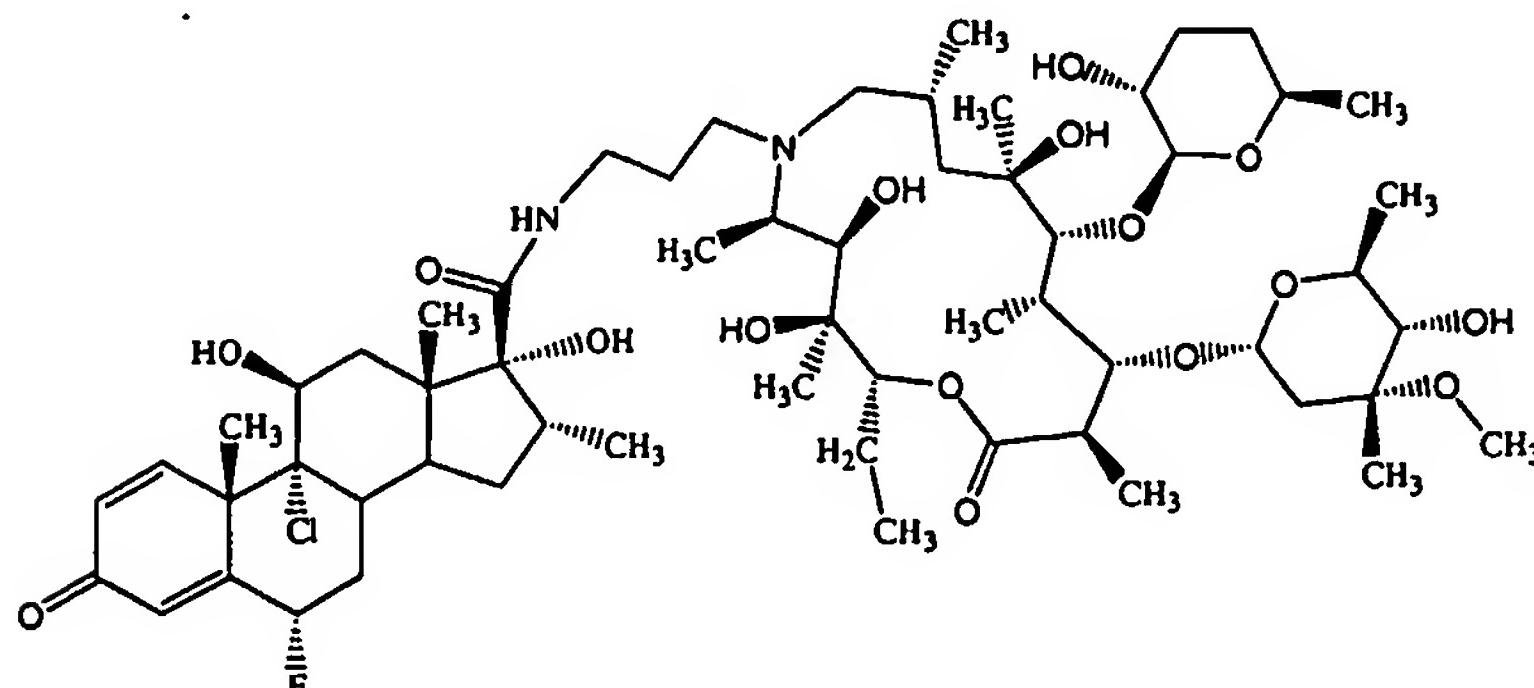
8. A compound 1 characterized by Formula:



9. A compound 2 characterized by Formula:

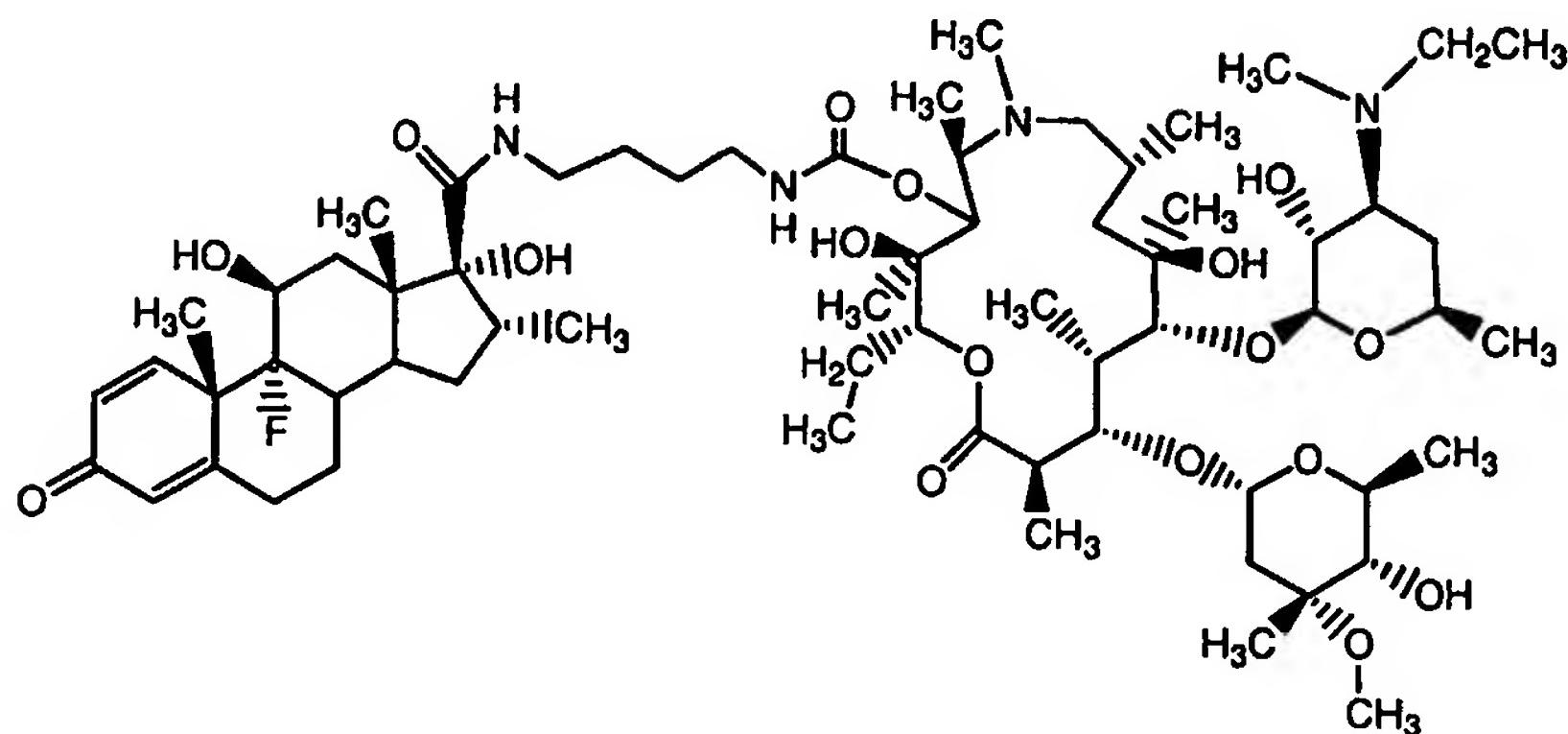


10. A compound 3 characterized by Formula:

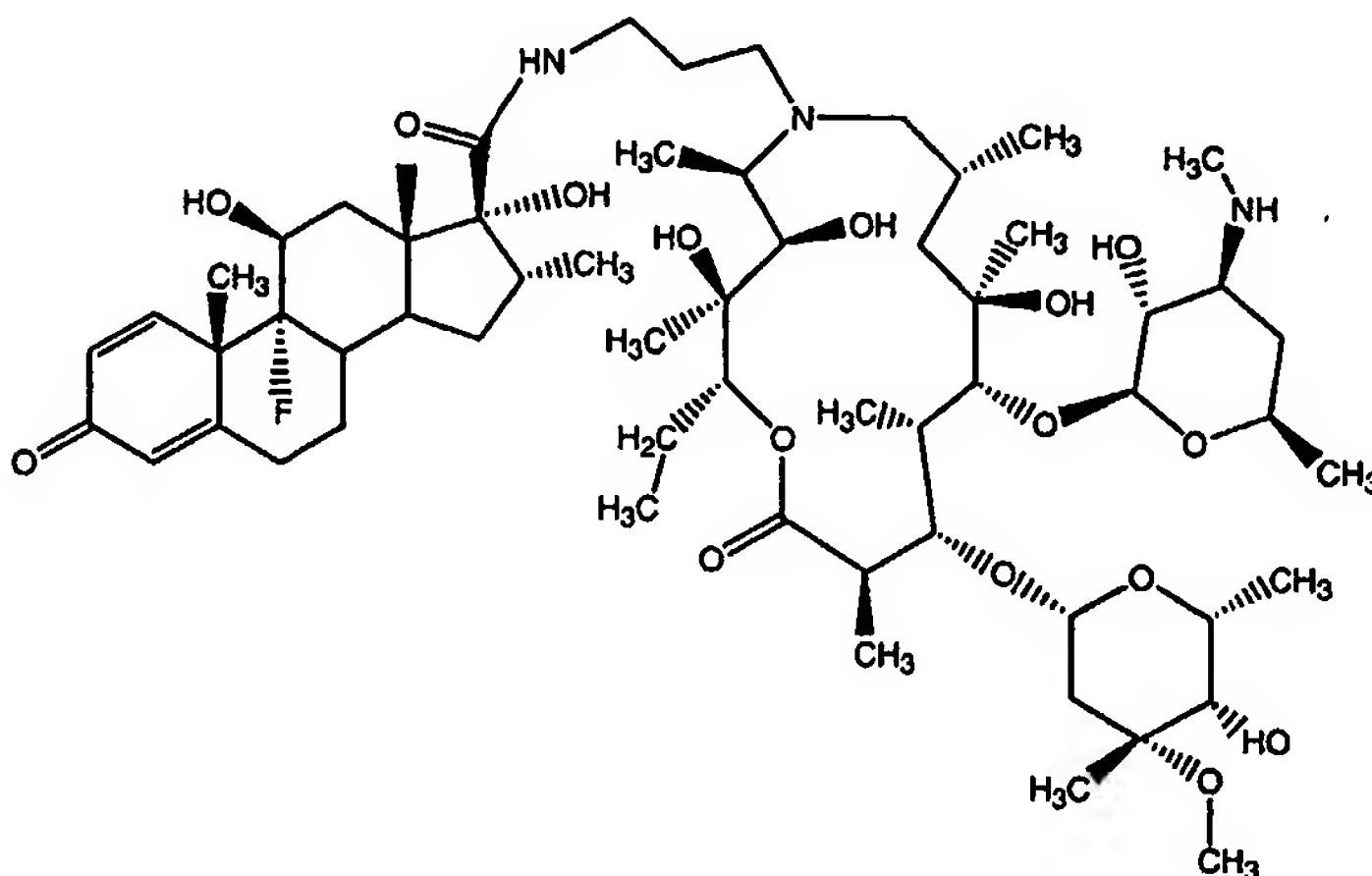


11. A compound 4 characterized by Formula:

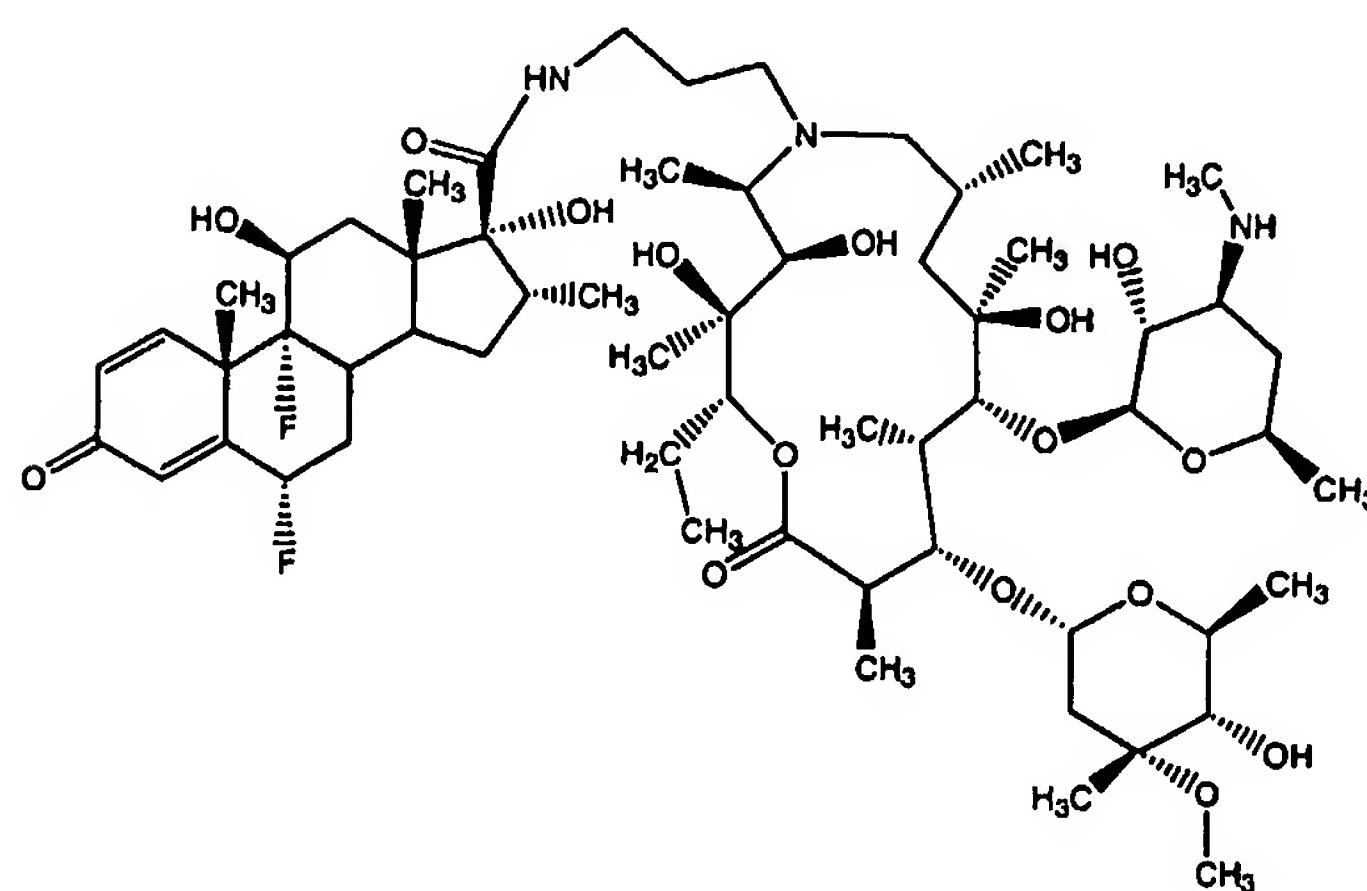
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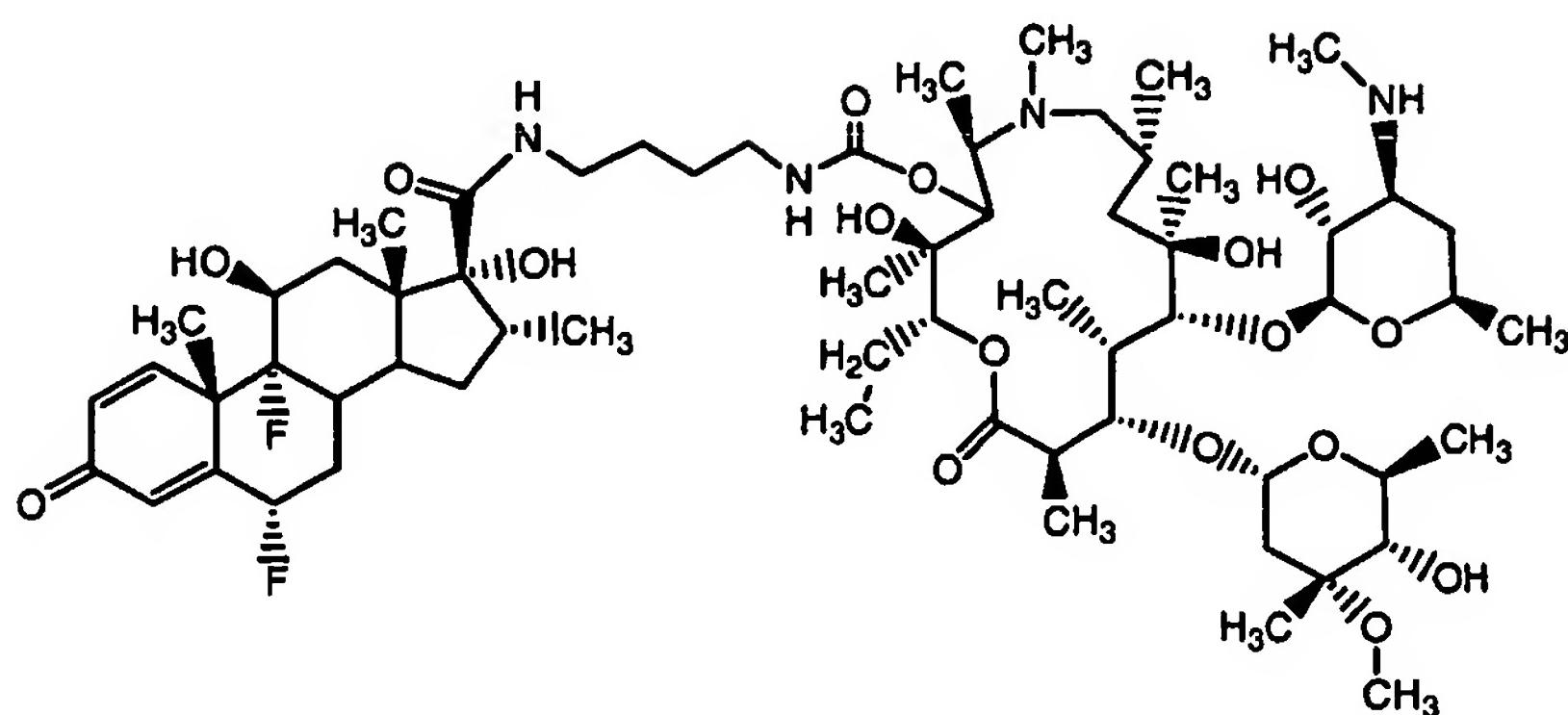
12. A compound 5 characterized by Formula:



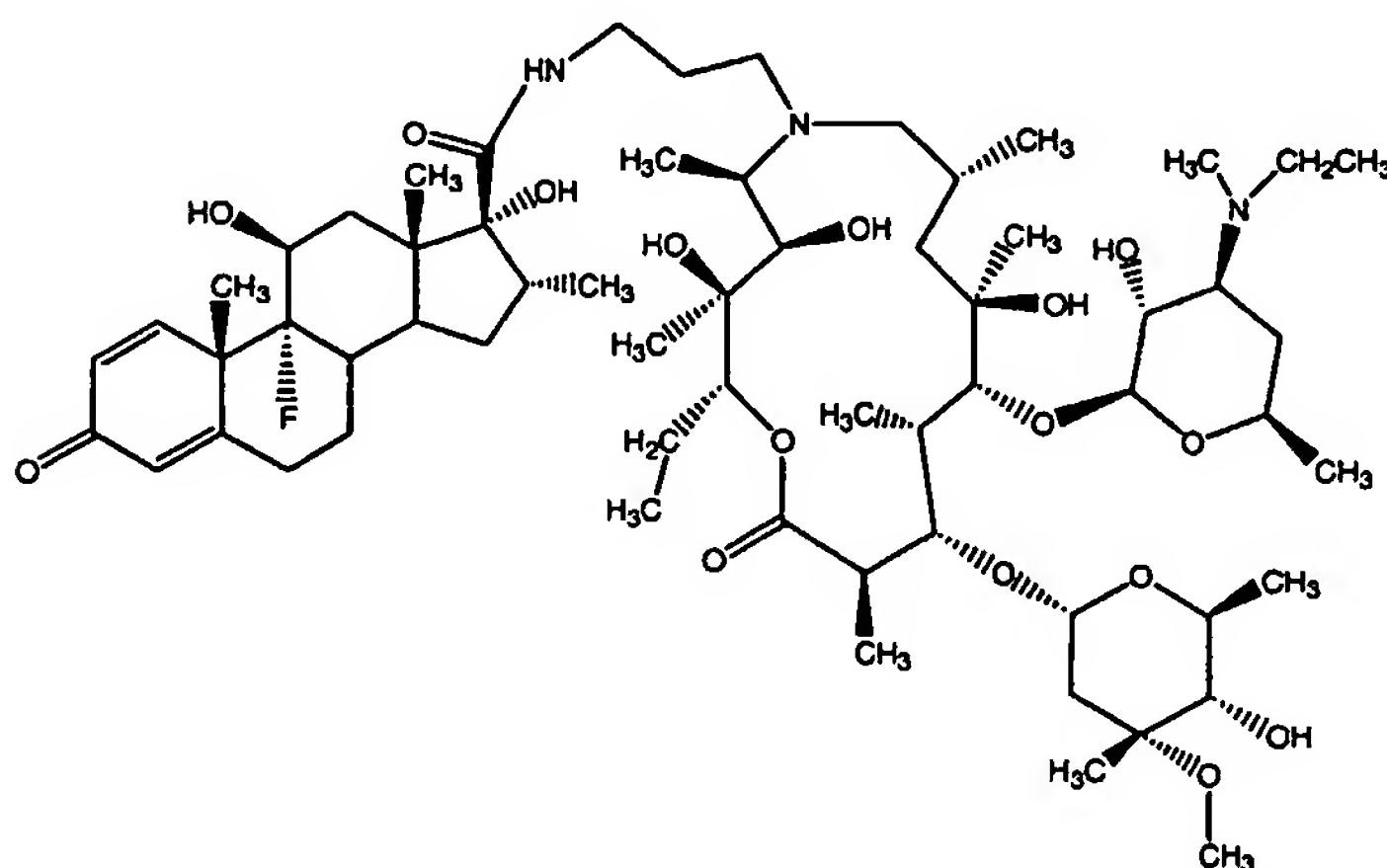
13. A compound 6 characterized by Formula:



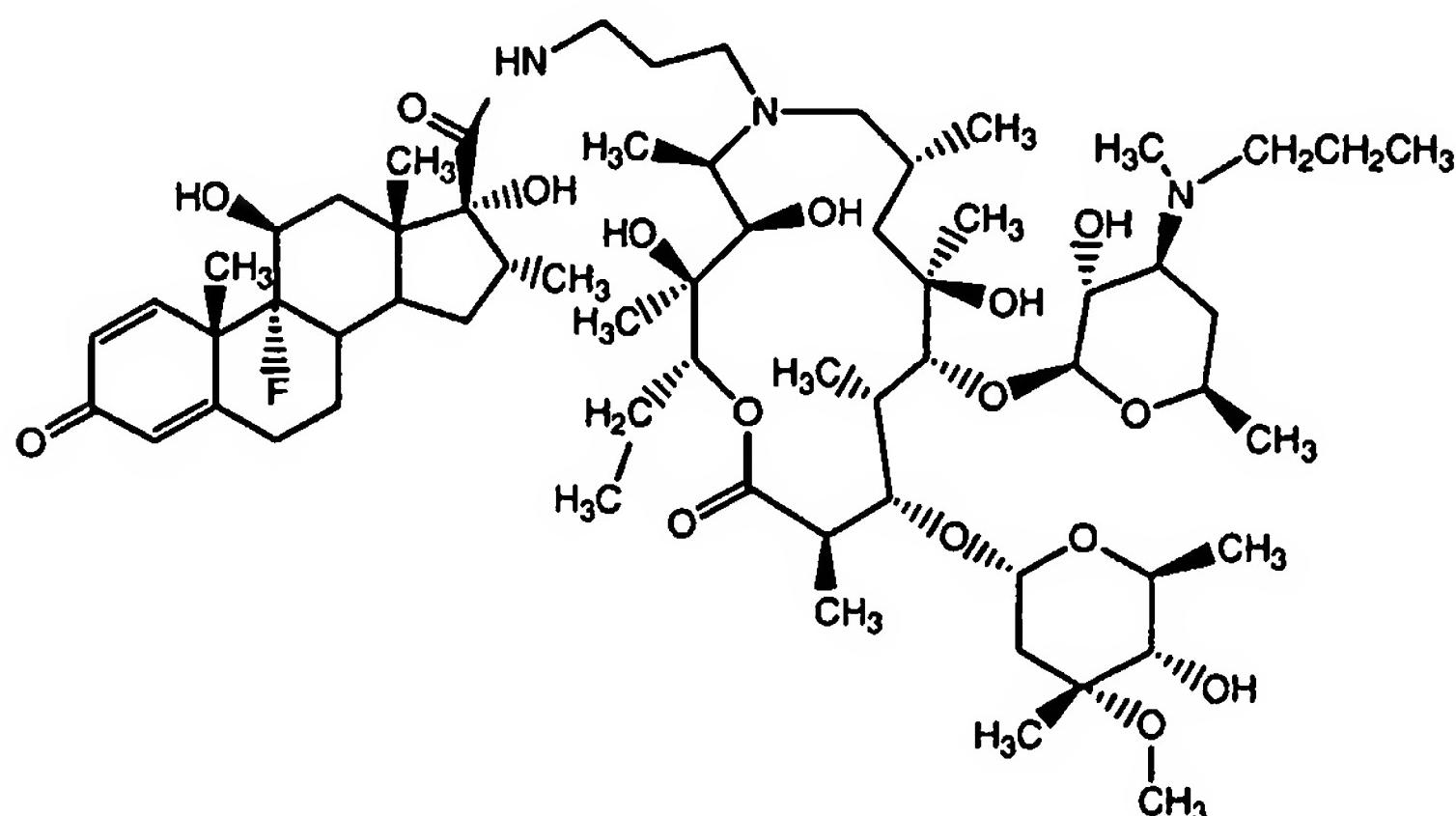
14. A compound 7 characterized by Formula:



15. A compound 8 characterized by Formula:

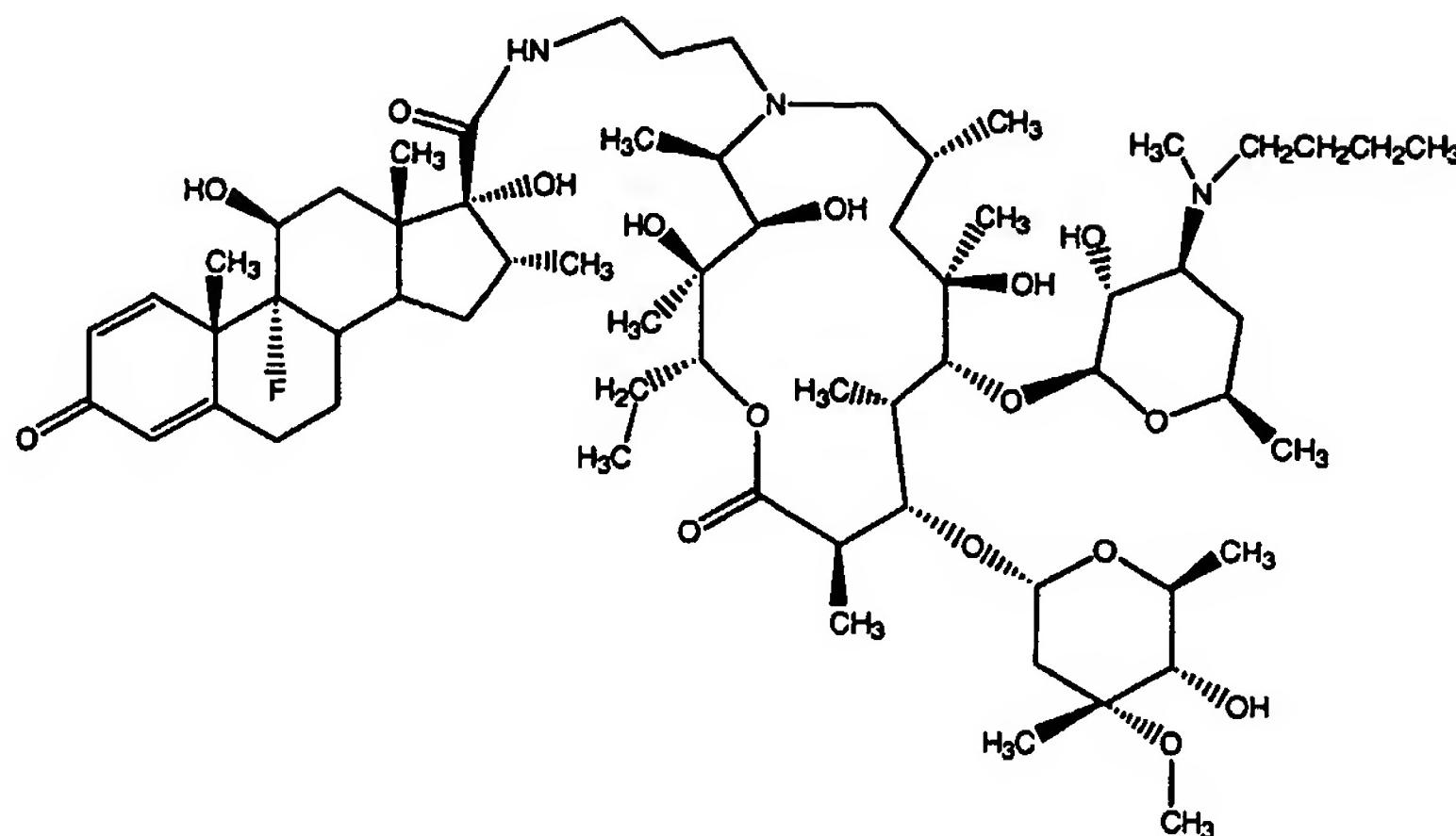


16. A compound 9 characterized by Formula:

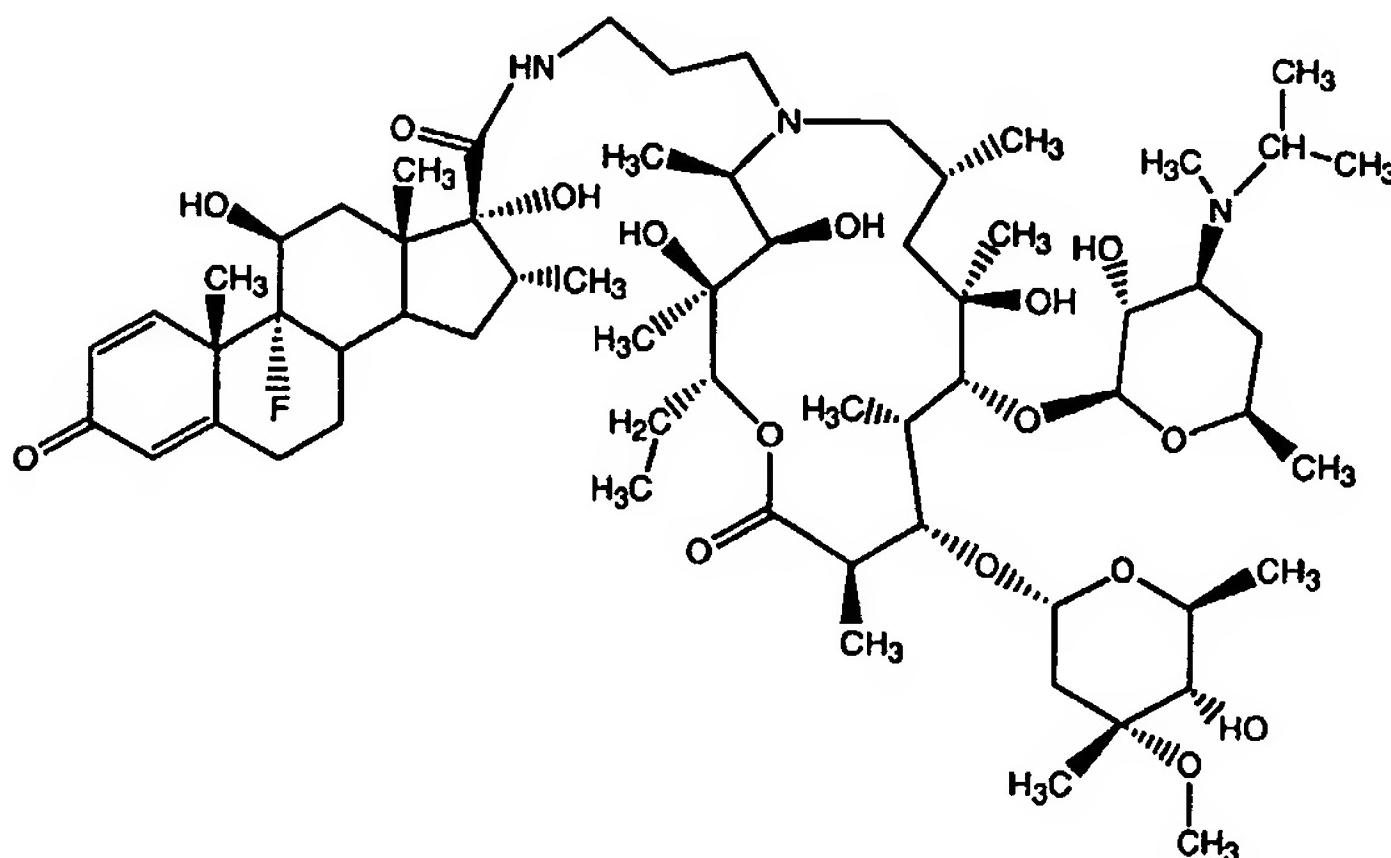


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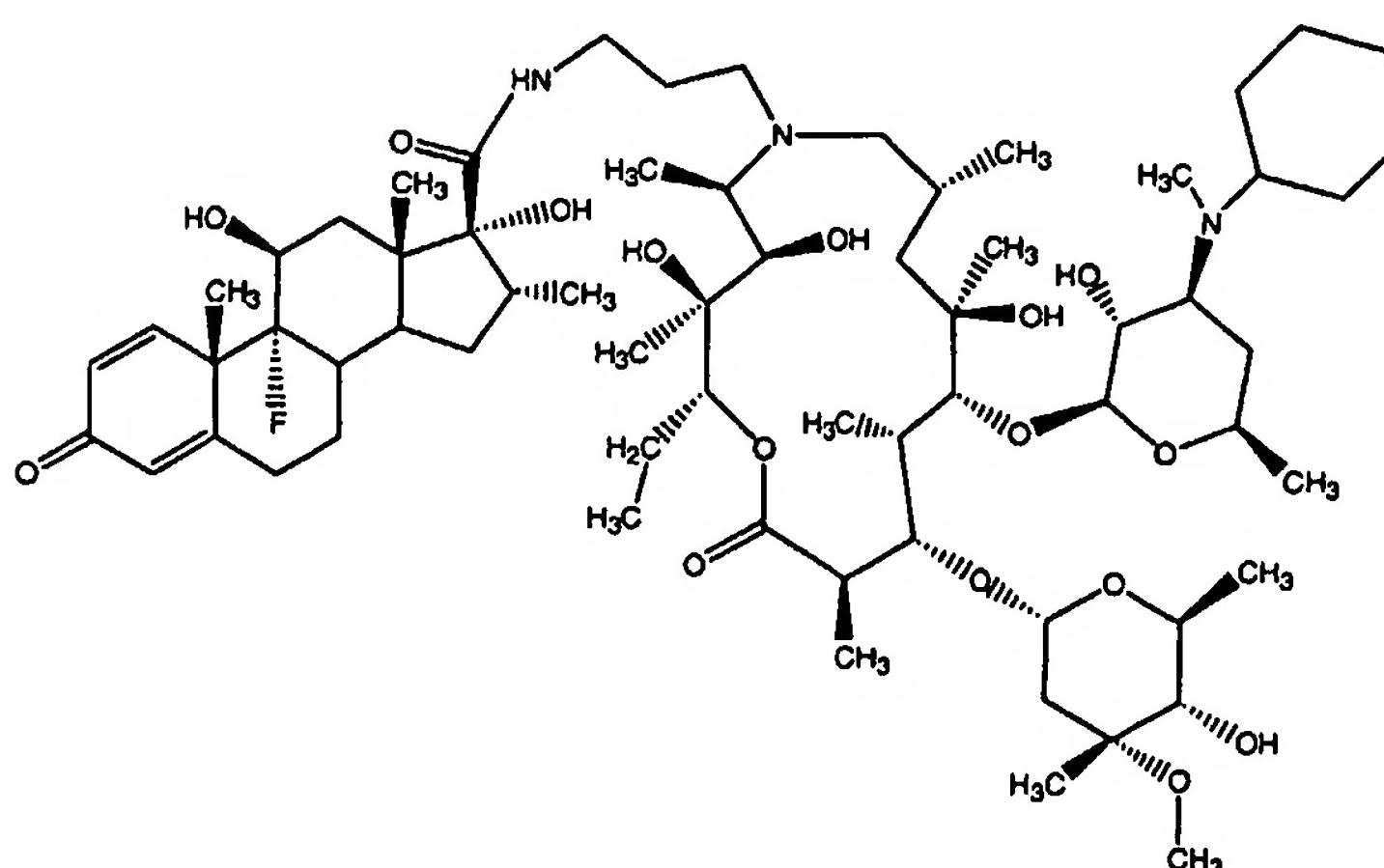
17. A compound 10 characterized by Formula:



18. A compound 11 characterized by Formula:

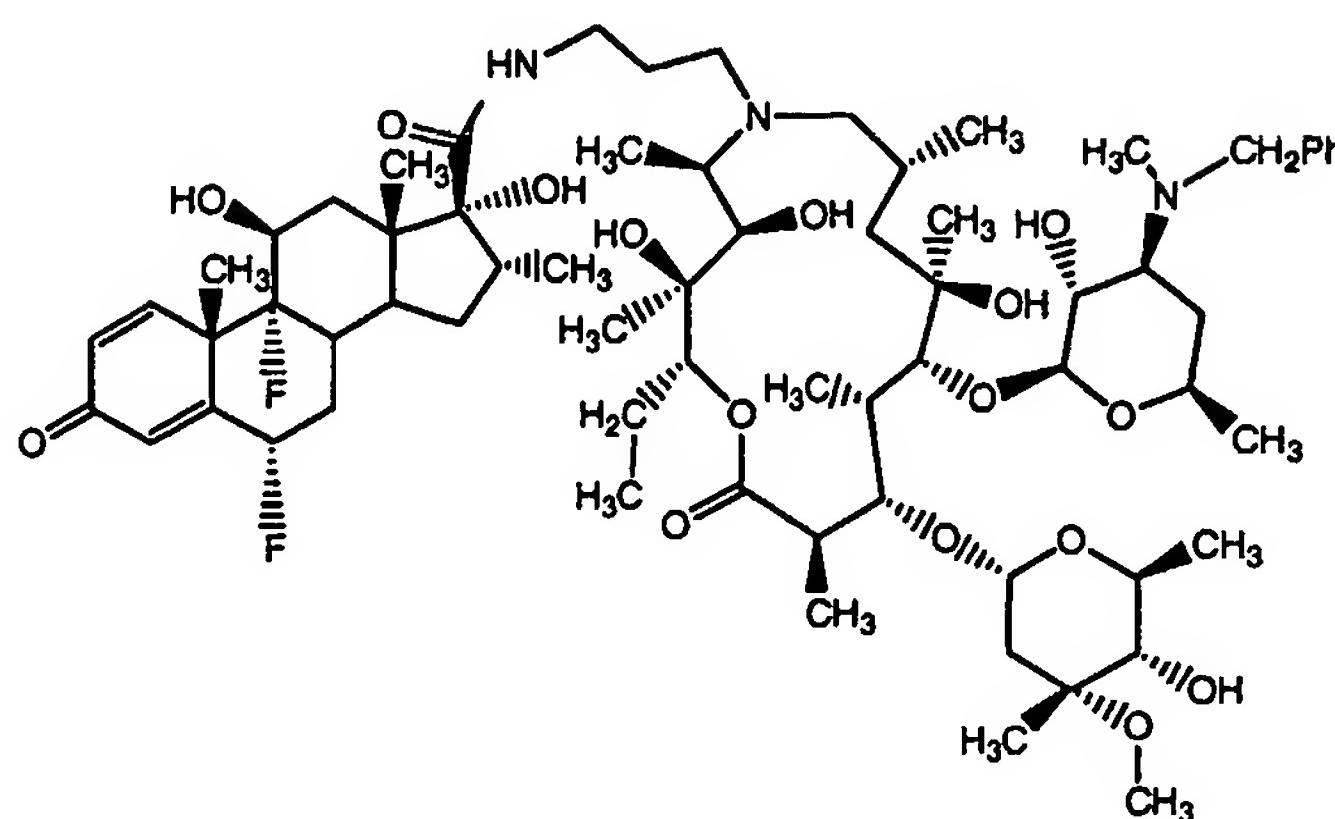


19. A compound 12 characterized by Formula:

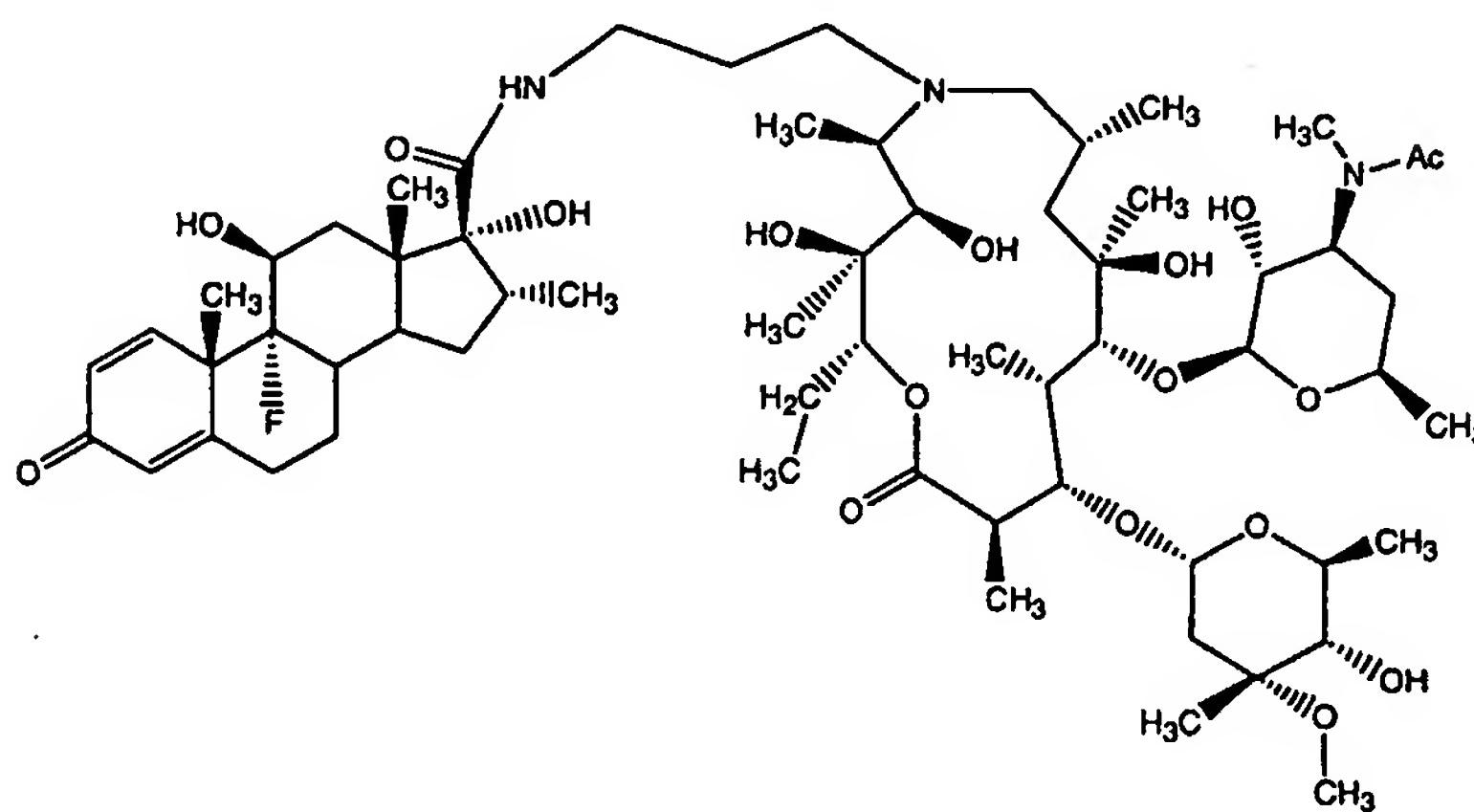


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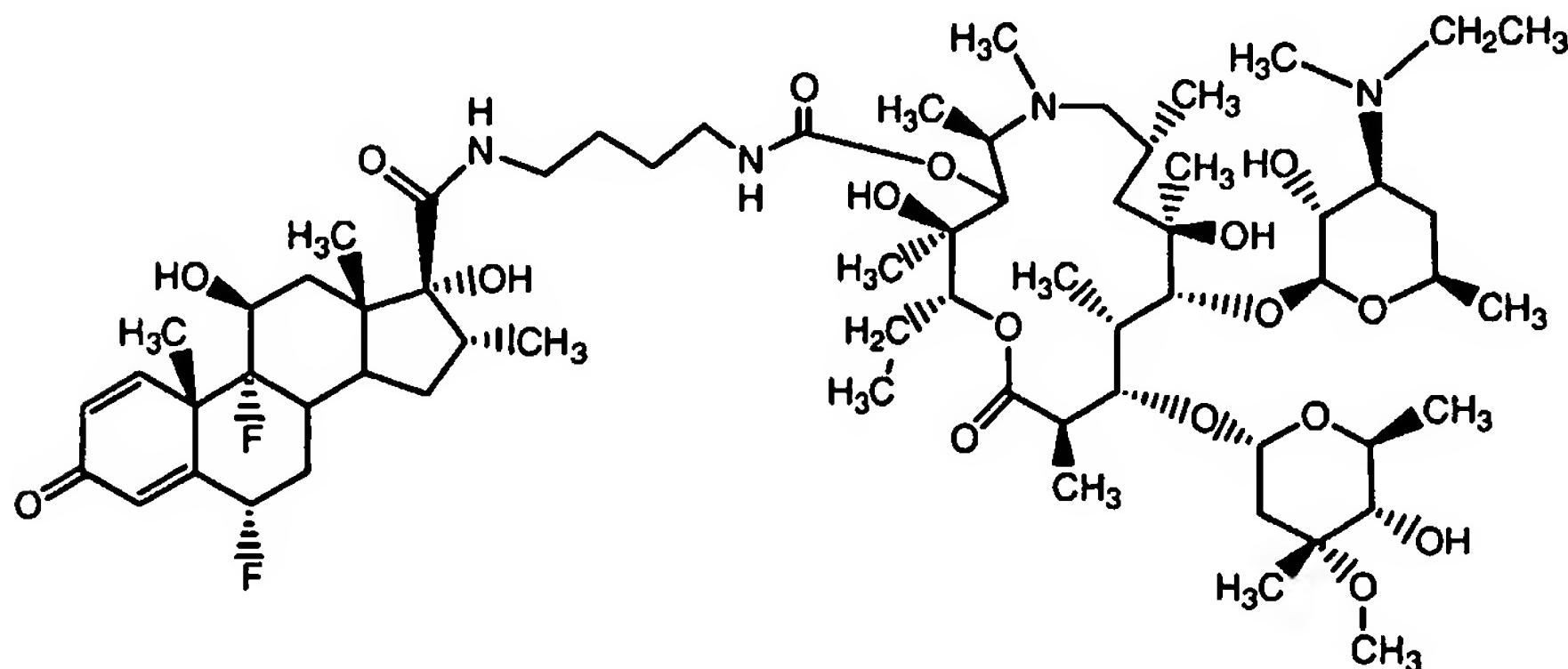
20. A compound 13 characterized by Formula:



21. A compound 14 characterized by Formula:

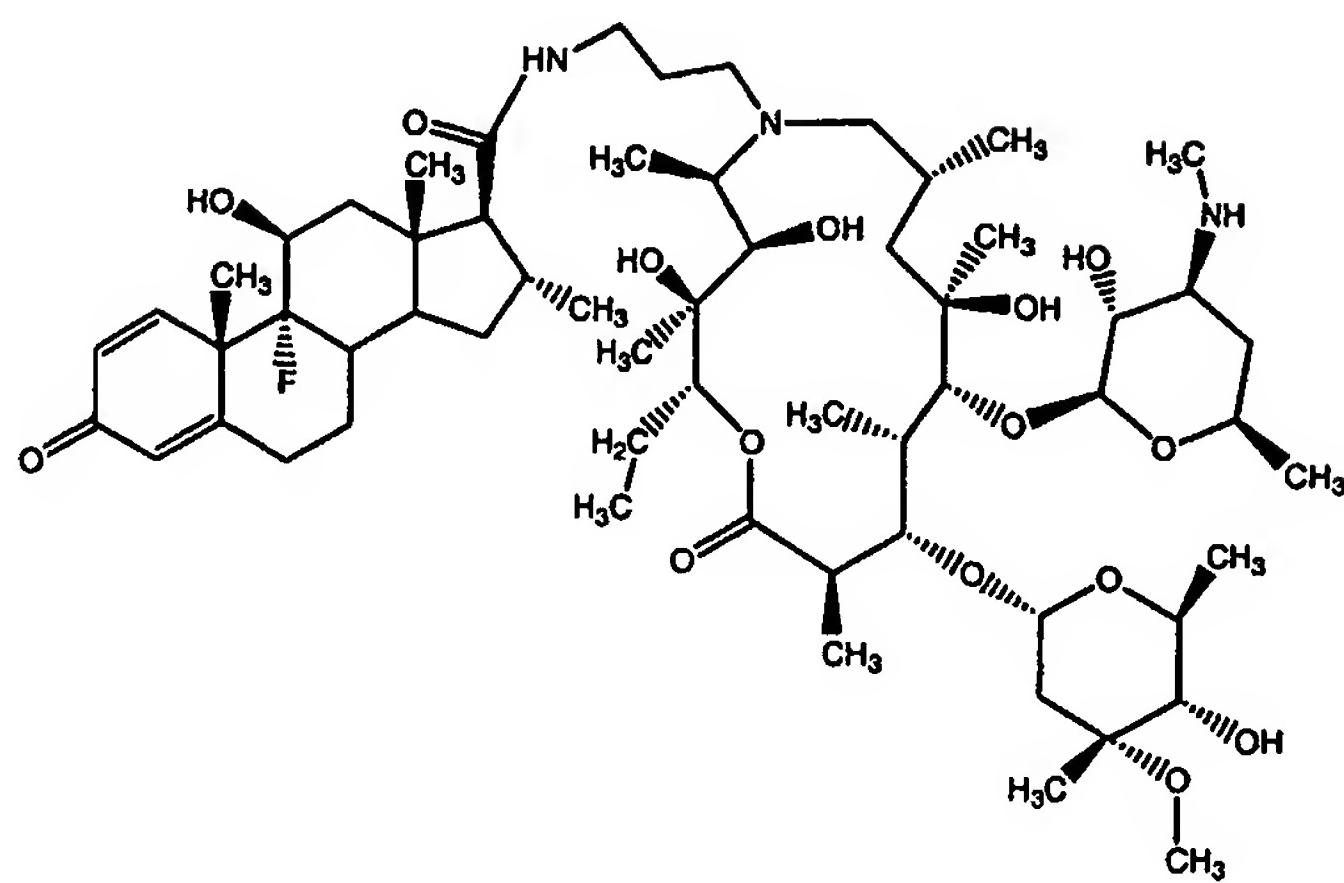


22. A compound 15 characterized by Formula:

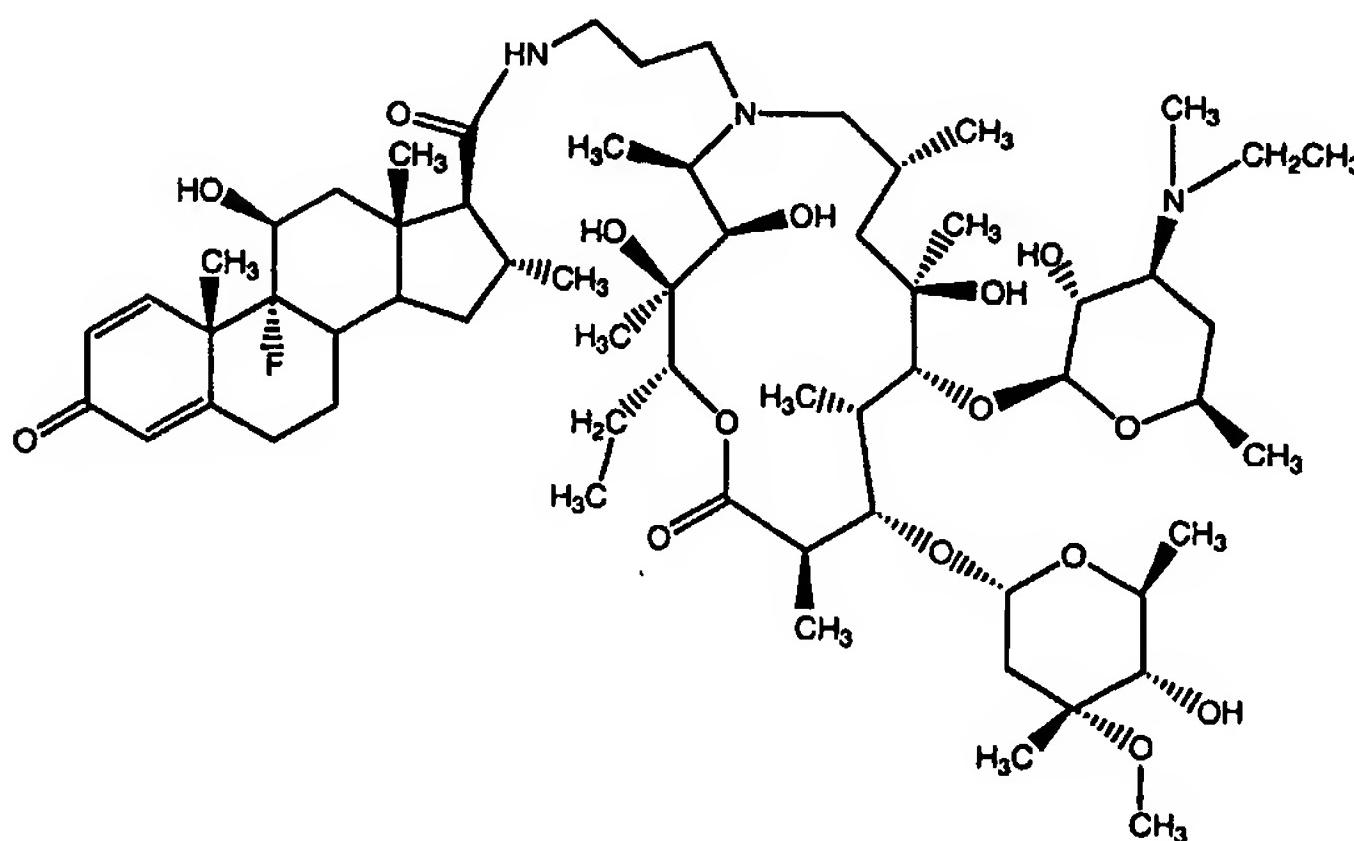


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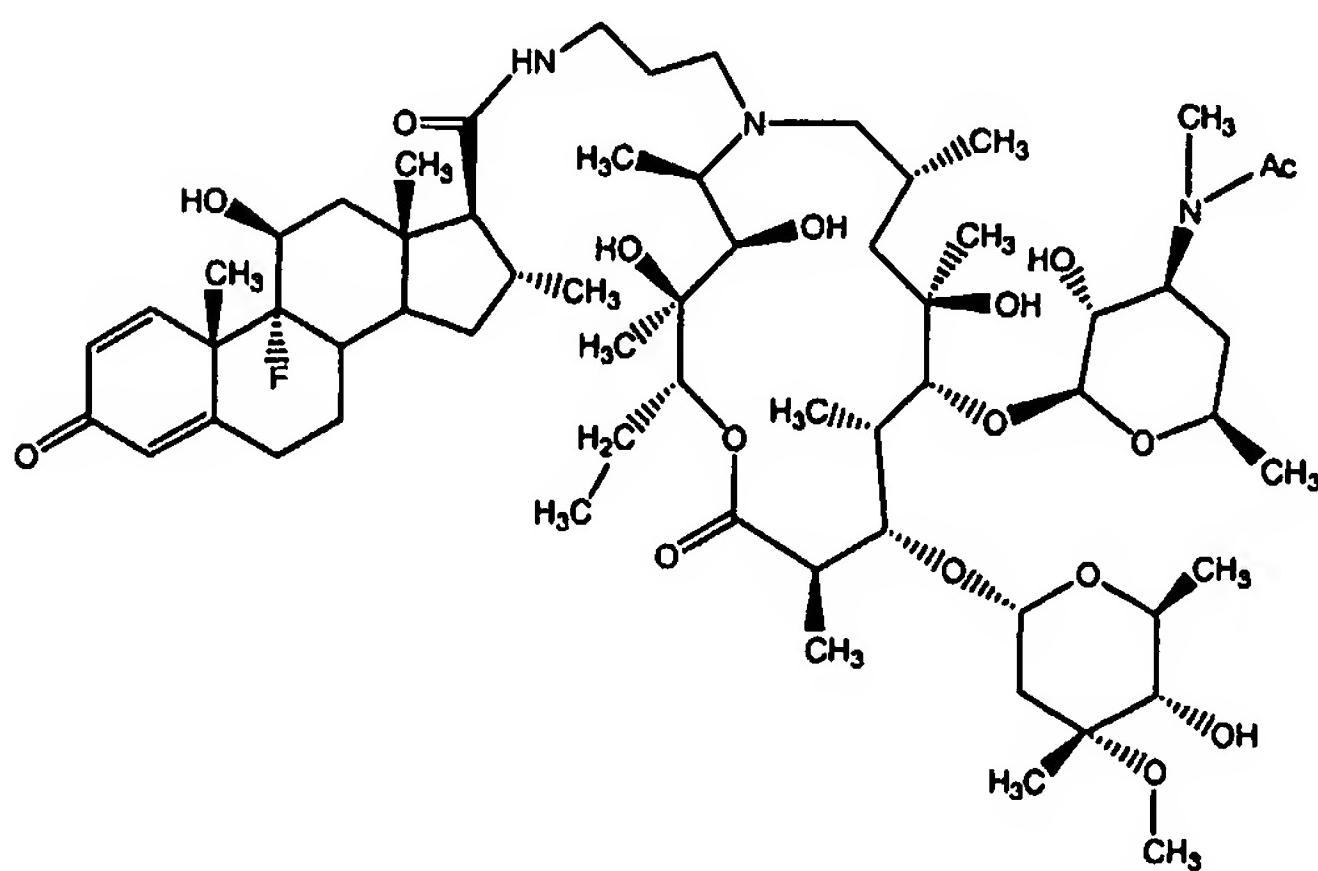
23. A compound 16 characterized by Formula:



24. A compound 17 characterized by Formula:

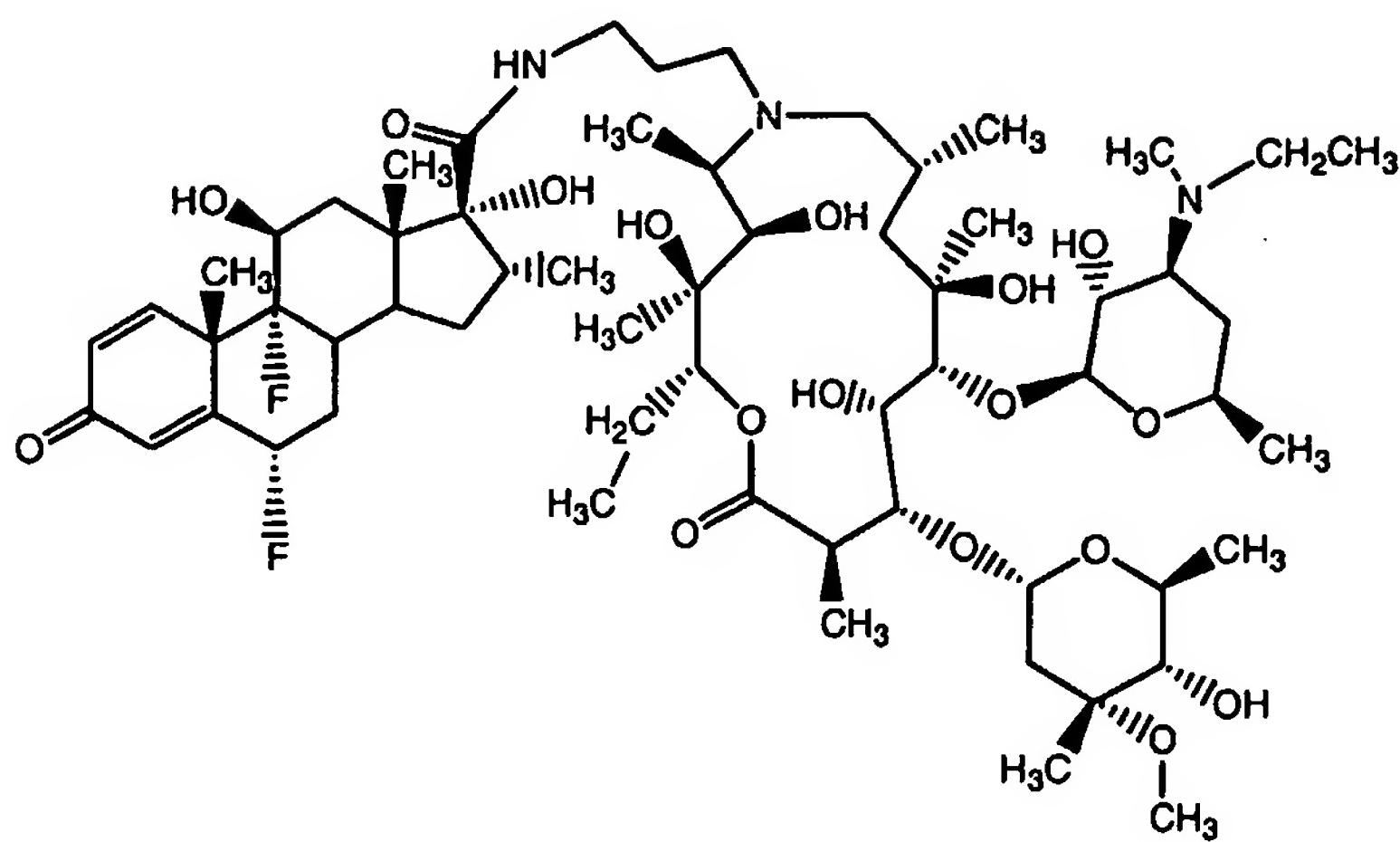


25. A compound 18 characterized by Formula:

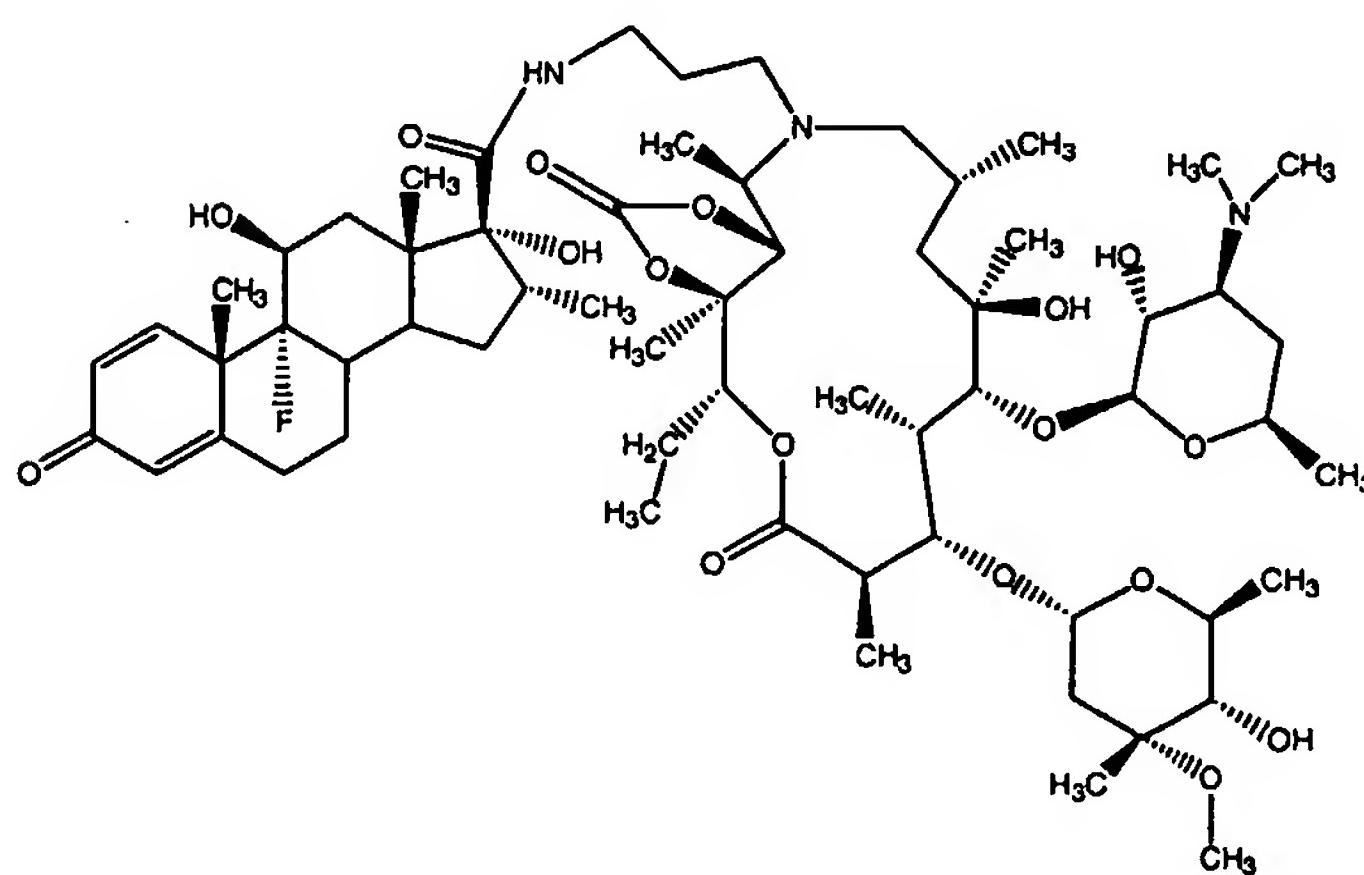


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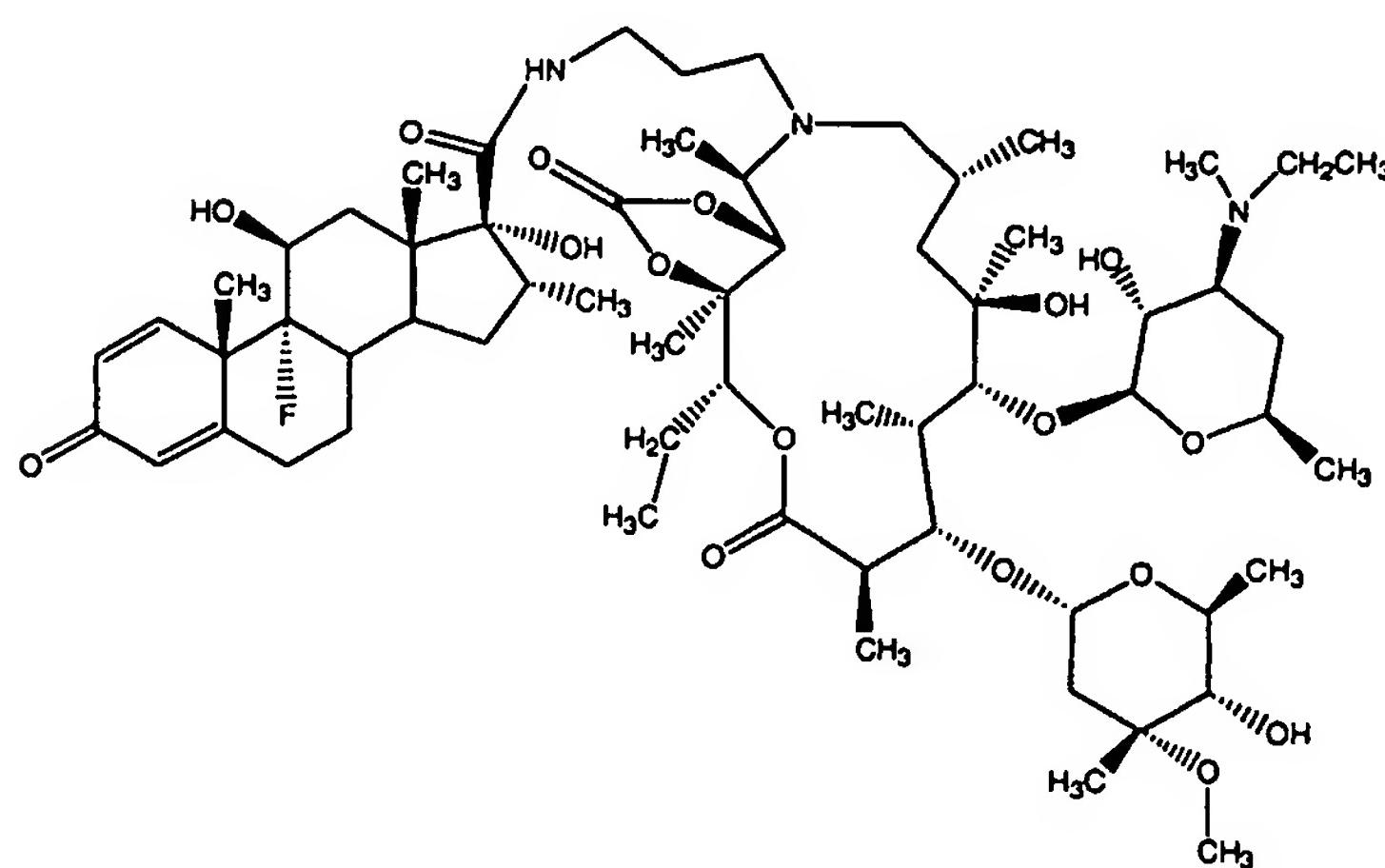
26. A compound 19 characterized by Formula:



27. A compound 20 characterized by Formula:

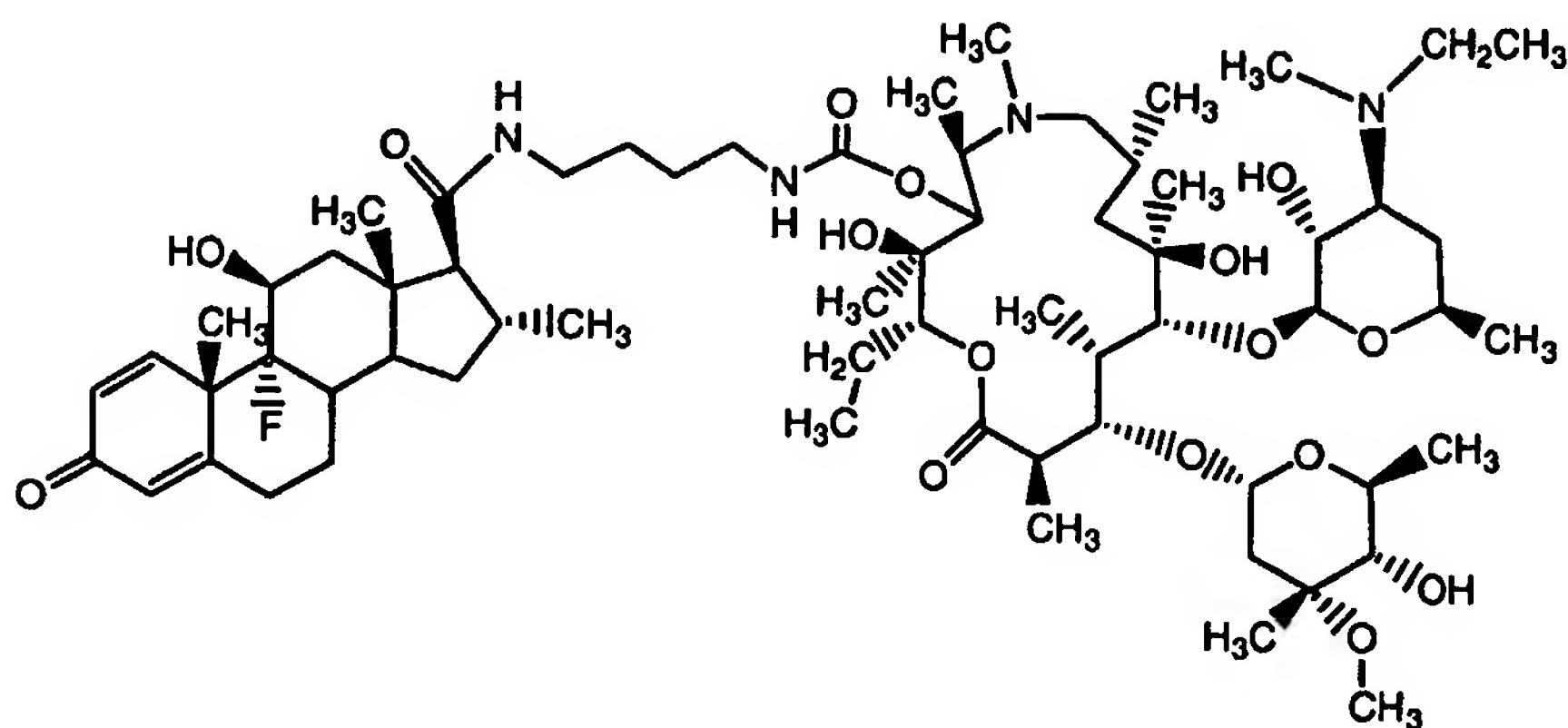


28. A compound 21 characterized by Formula:

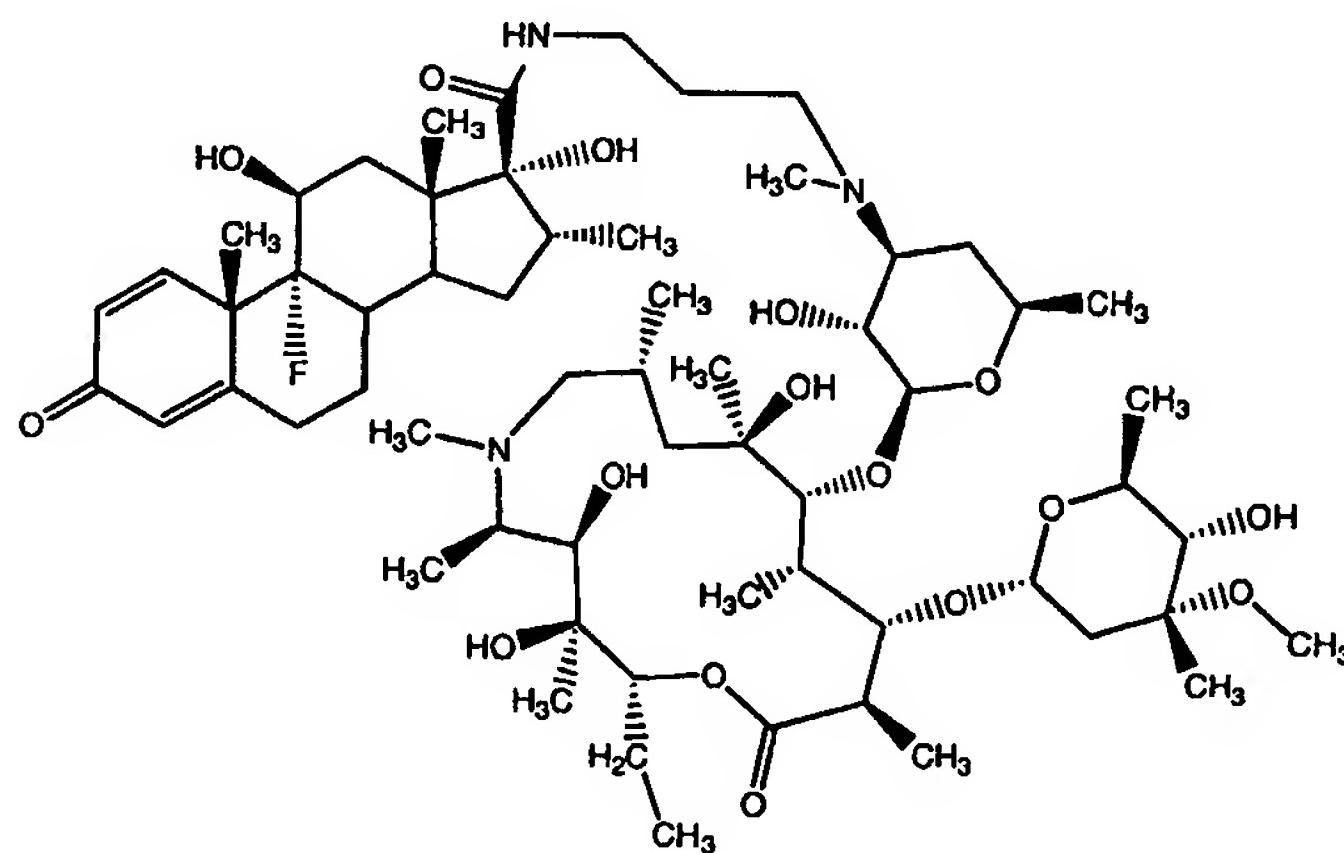


105

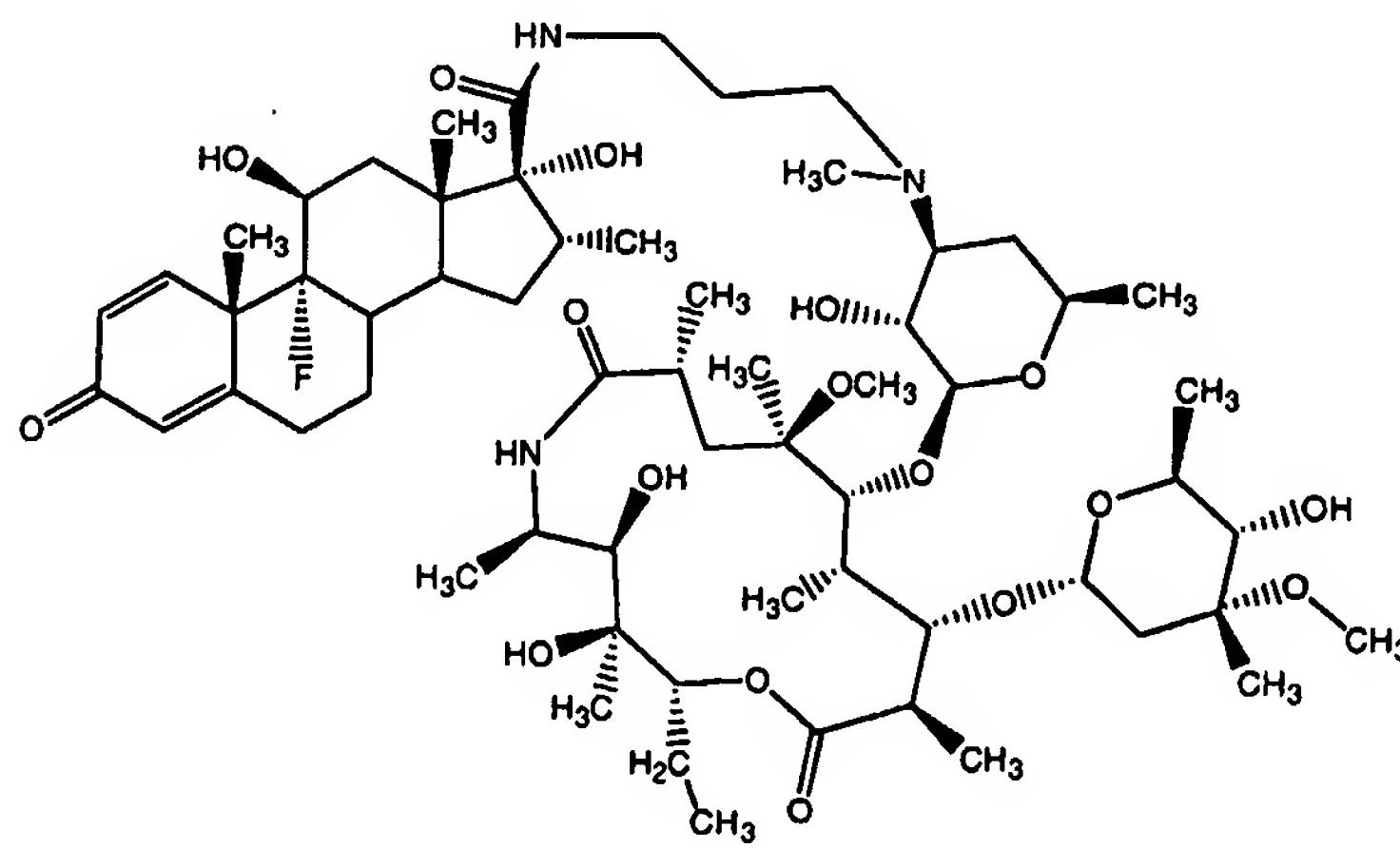
29. A compound 22 characterized by Formula:



30. A compound 23 characterized by Formula:

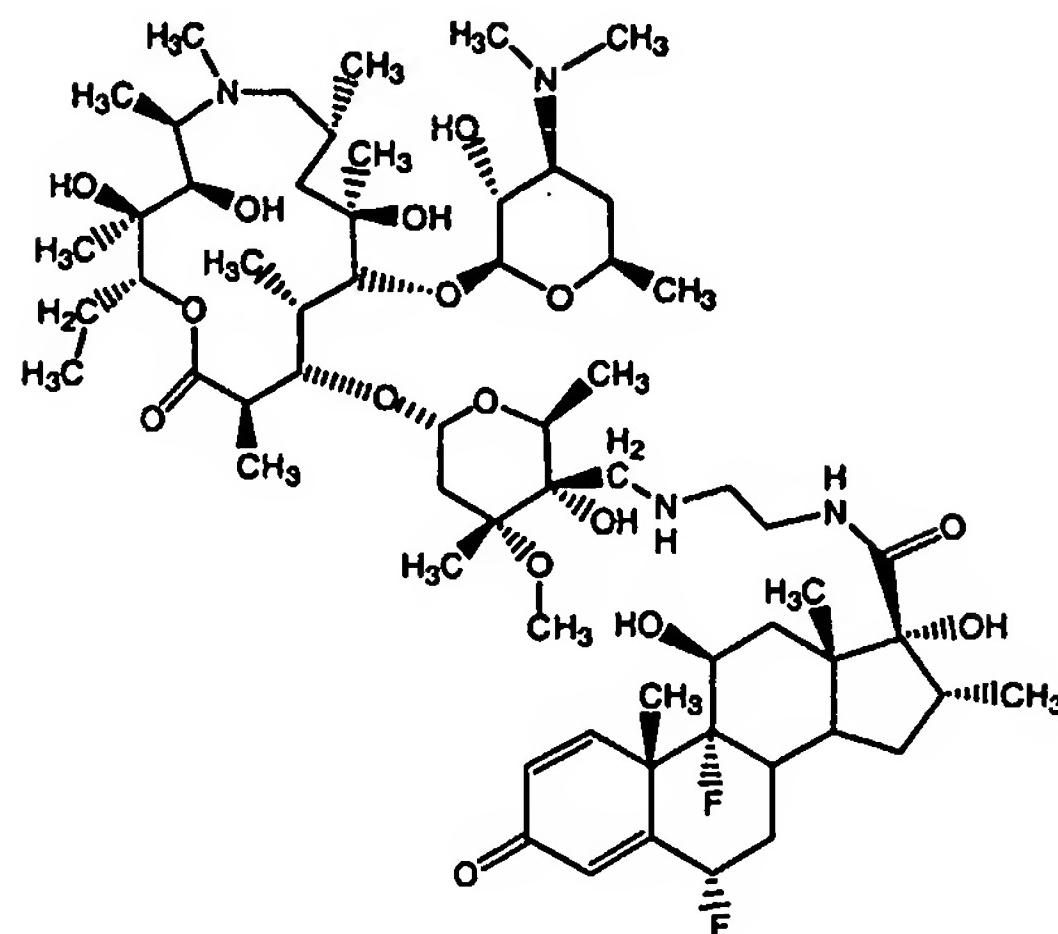


31. A compound 24 characterized by Formula:

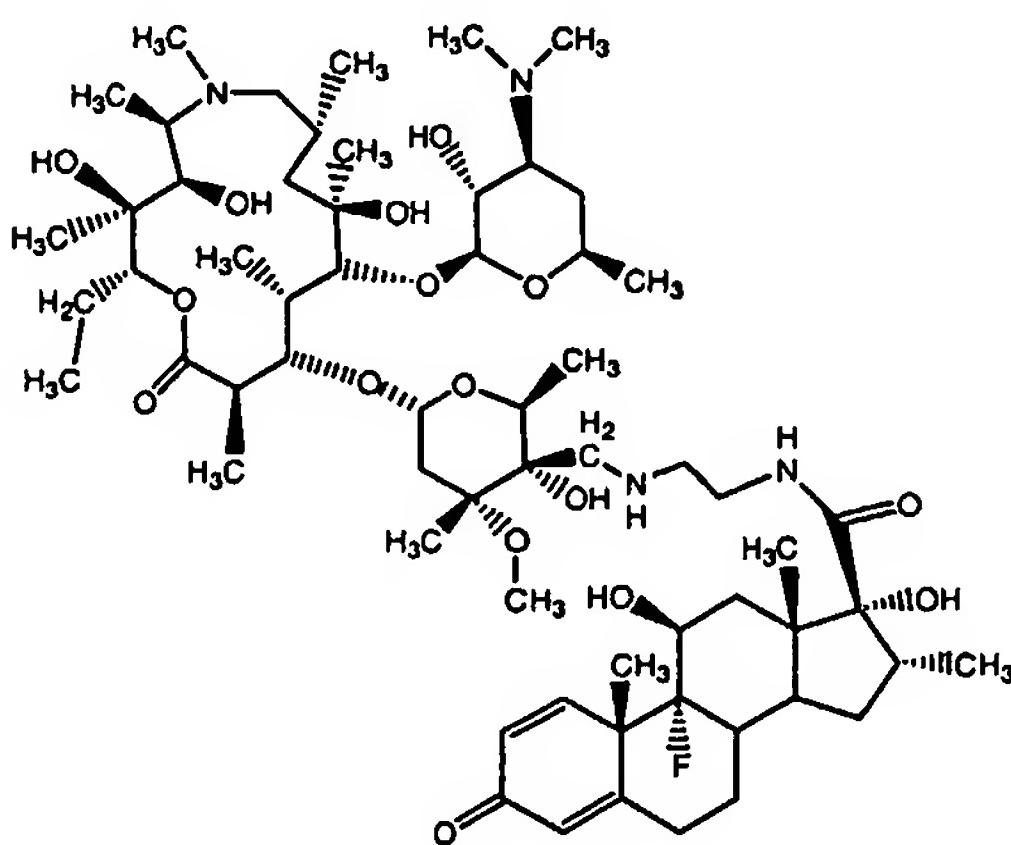


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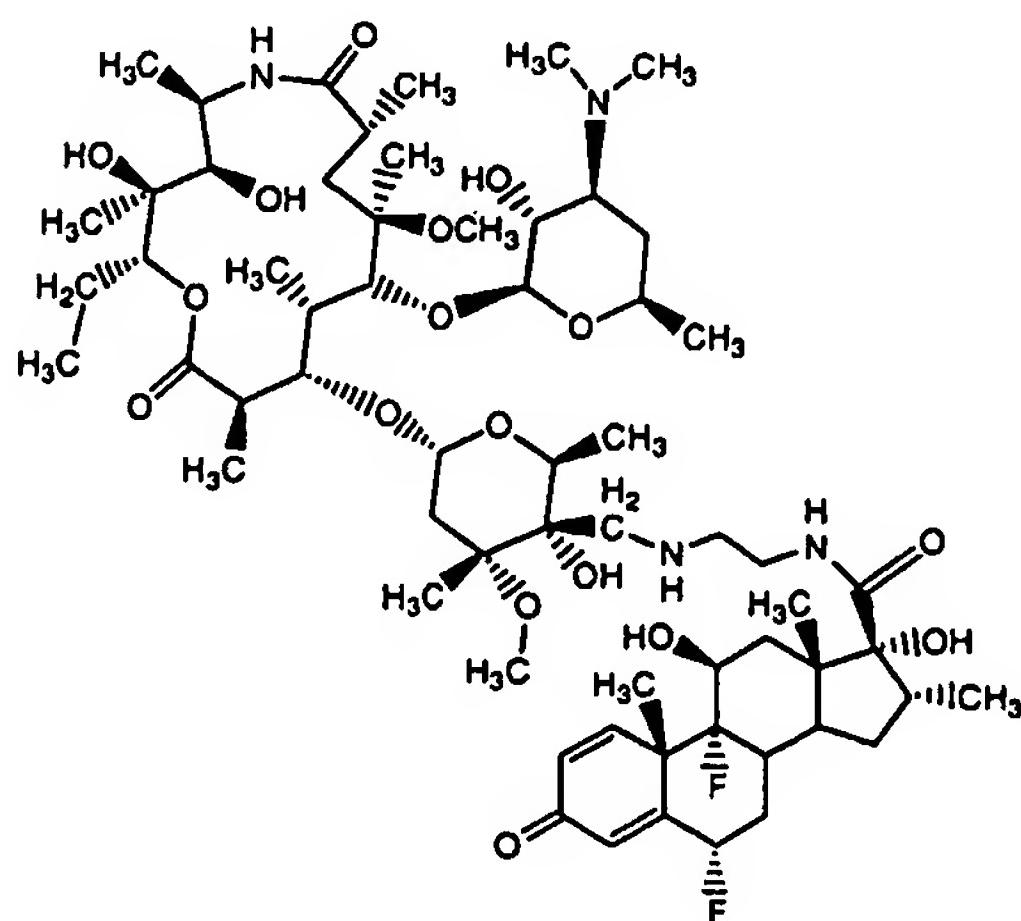
32. A compound 25 characterized by Formula:



33. A compound 26 characterized by Formula:

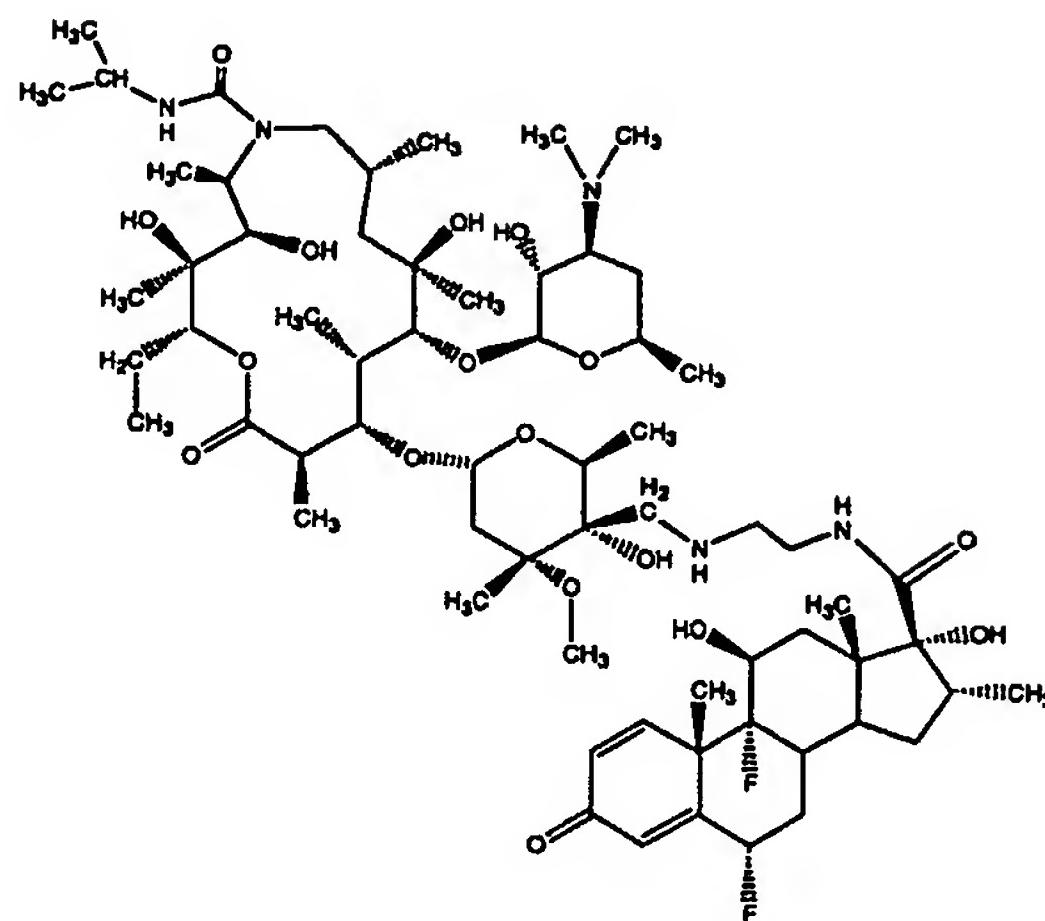


34. A compound 27 characterized by Formula:

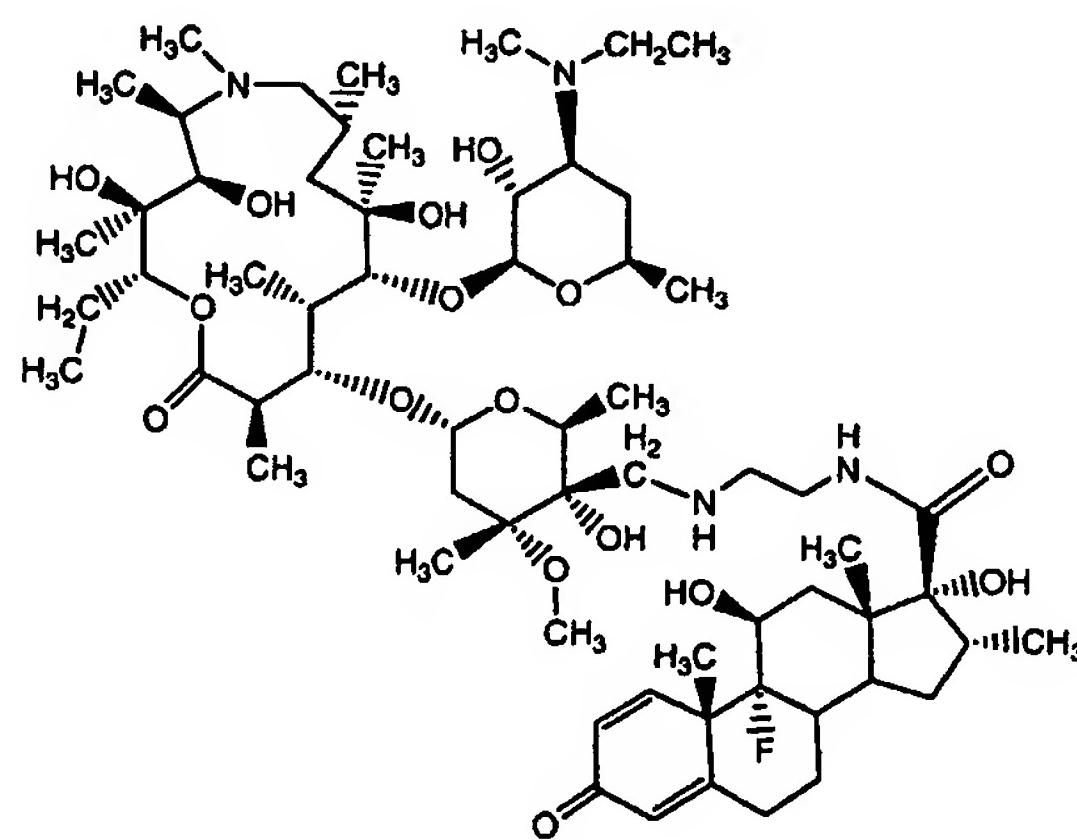


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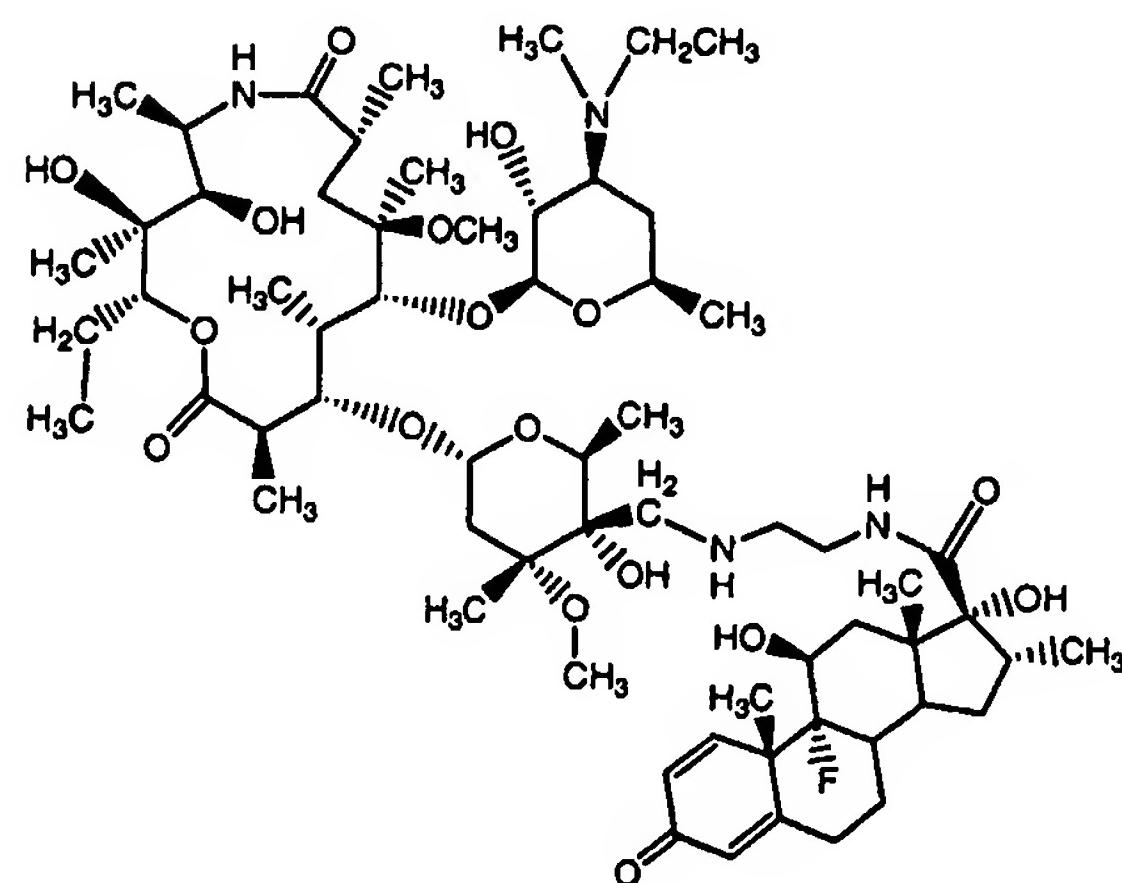
35. A compound 28 characterized by Formula:



36. A compound 29 characterized by Formula:

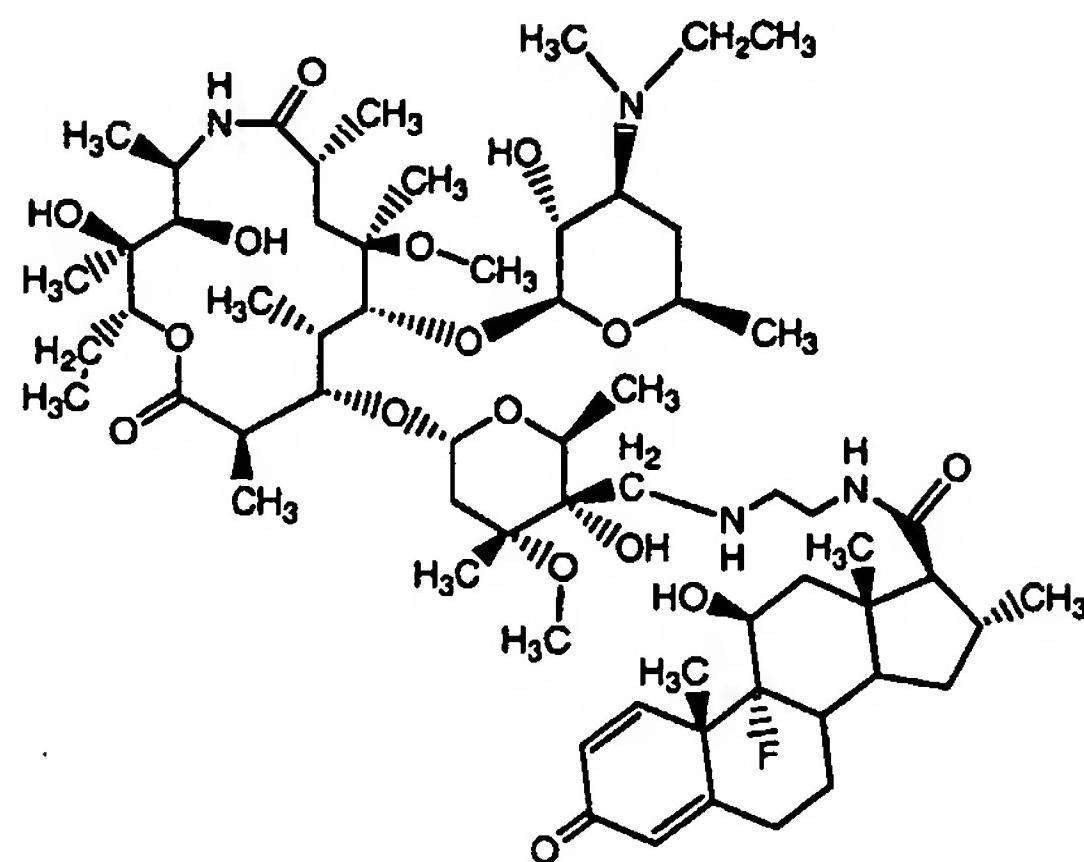


37. A compound 30 characterized by Formula:

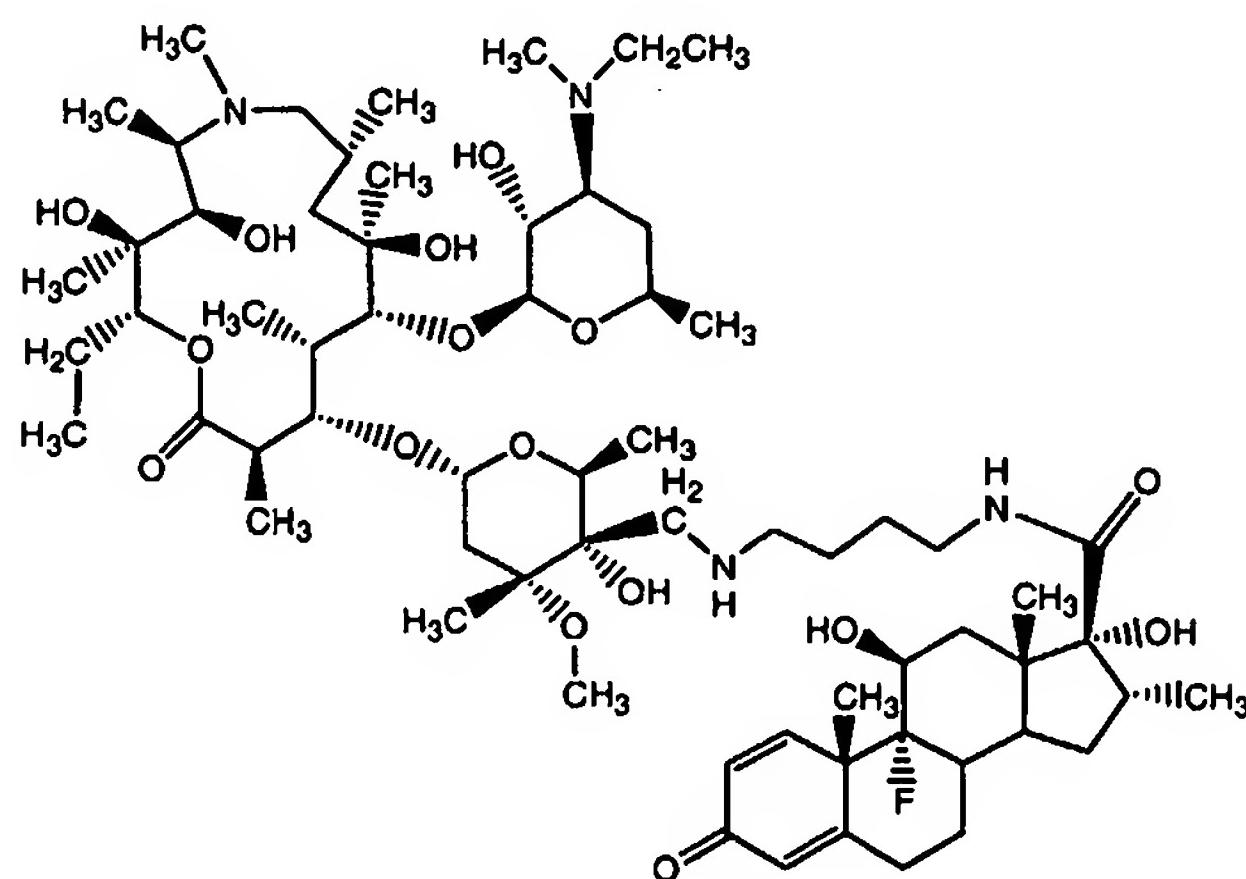


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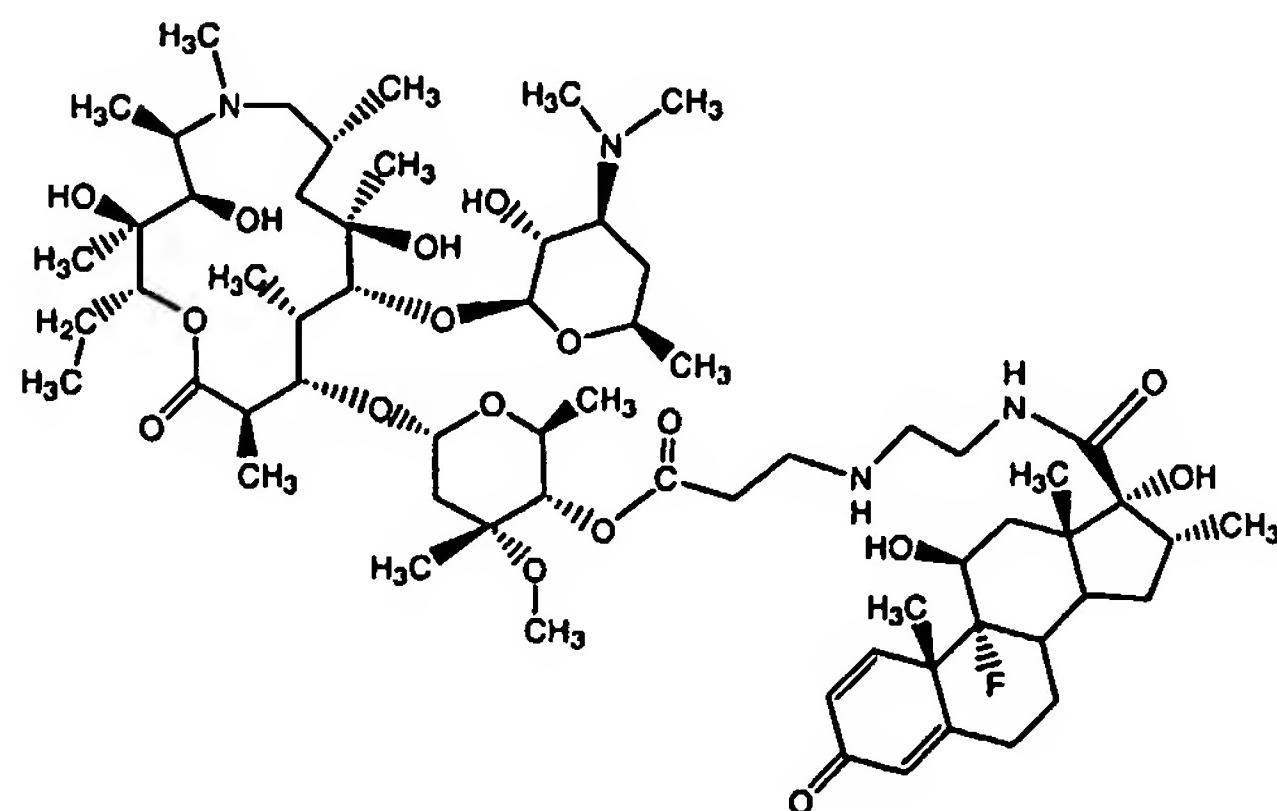
38. A compound 31 characterized by Formula:



39. A compound 32 characterized by Formula:

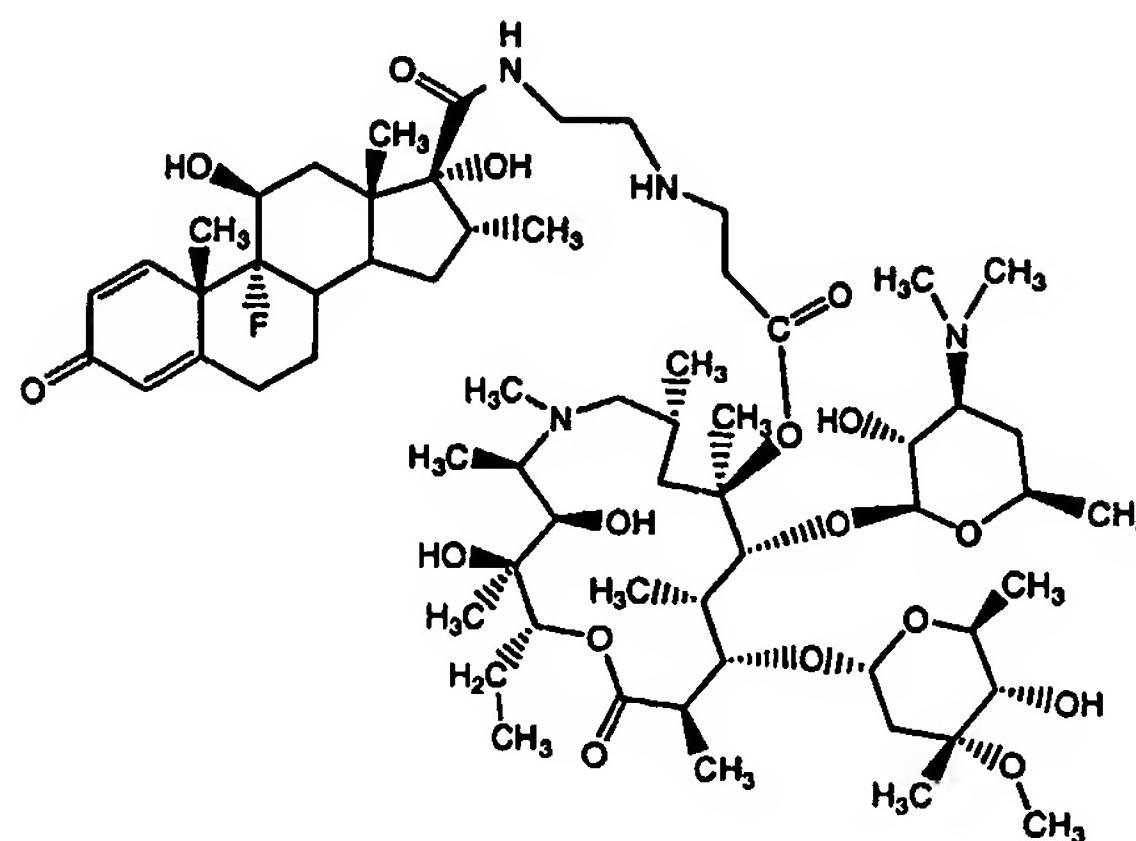


40. A compound 33 characterized by Formula:

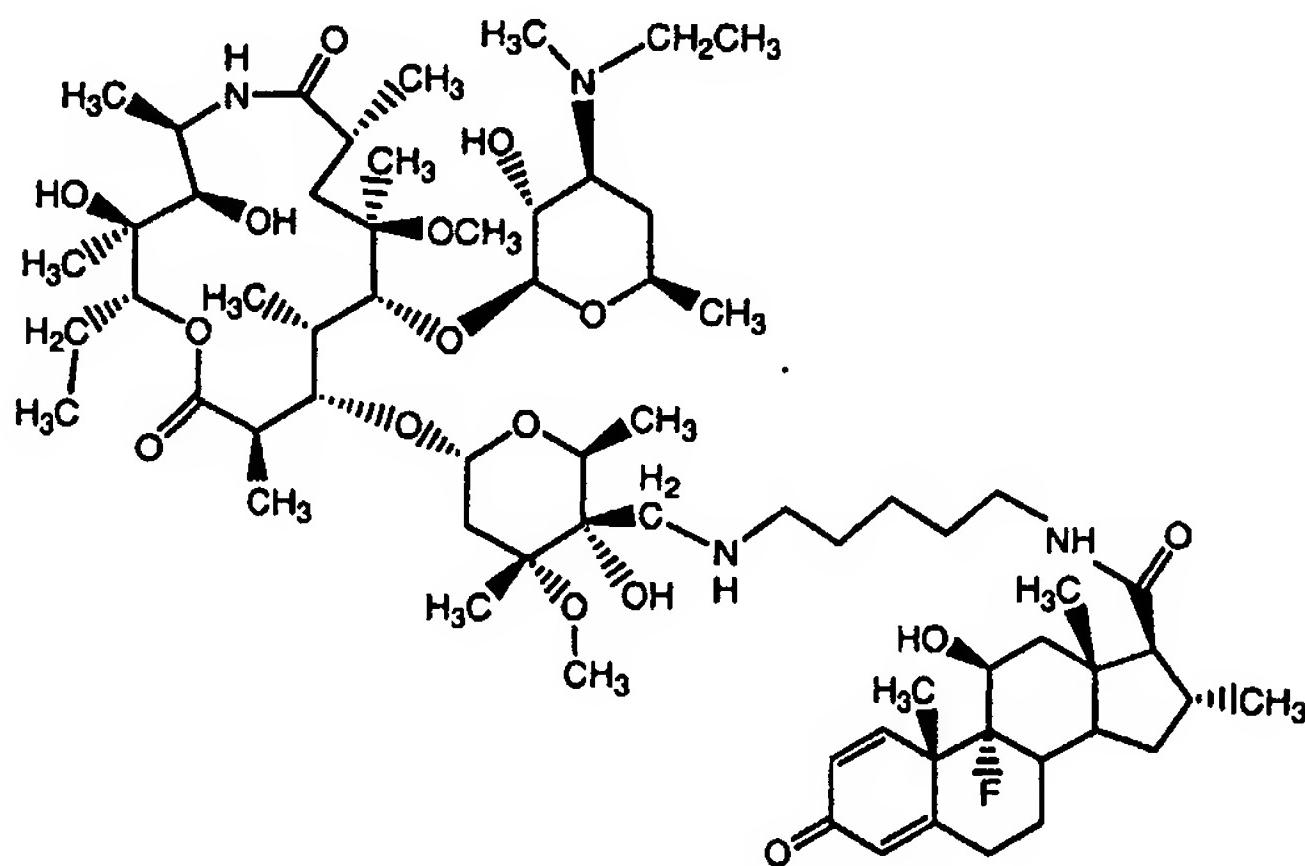


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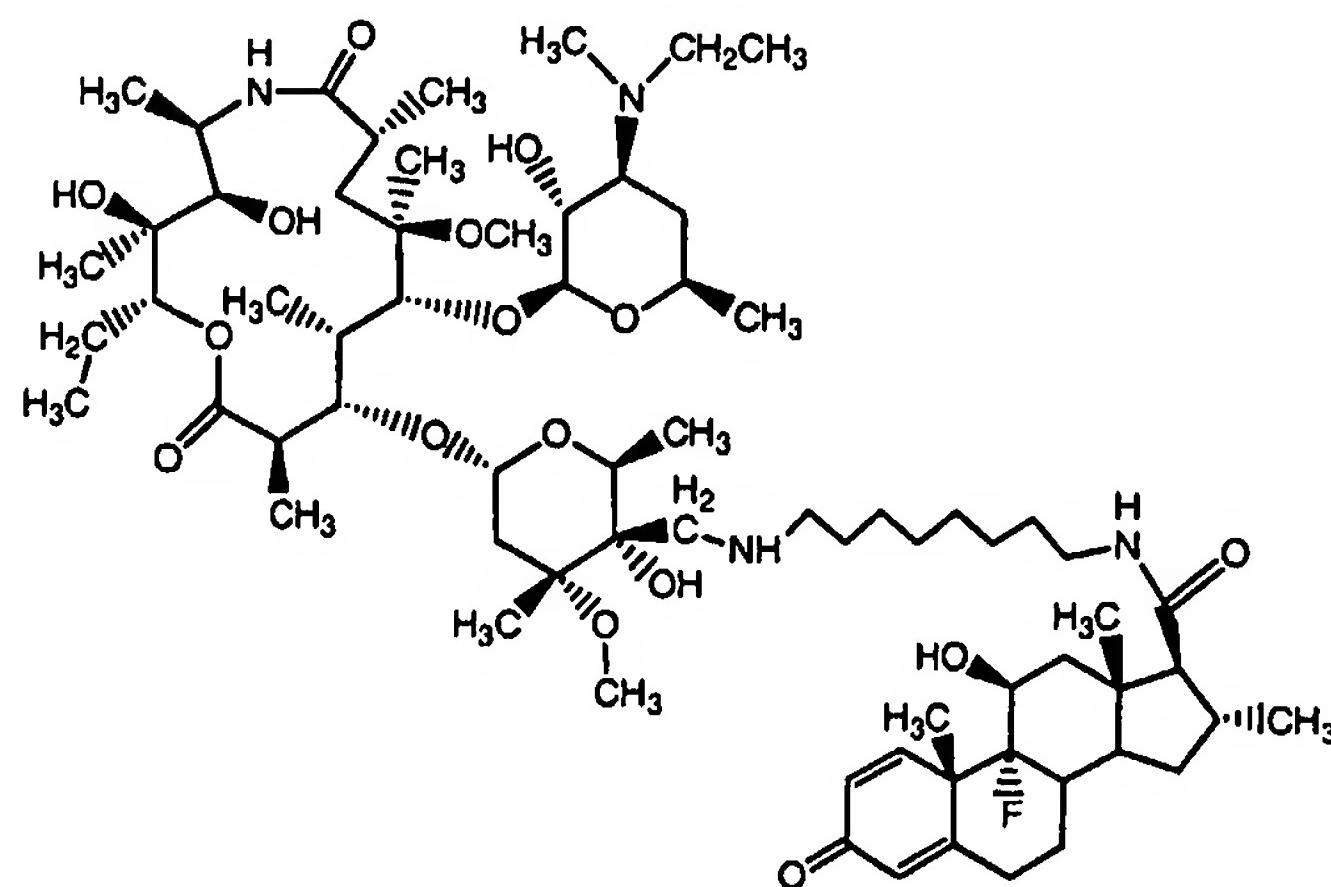
41. A compound 34 characterized by Formula:



42. A compound 35 characterized by Formula:

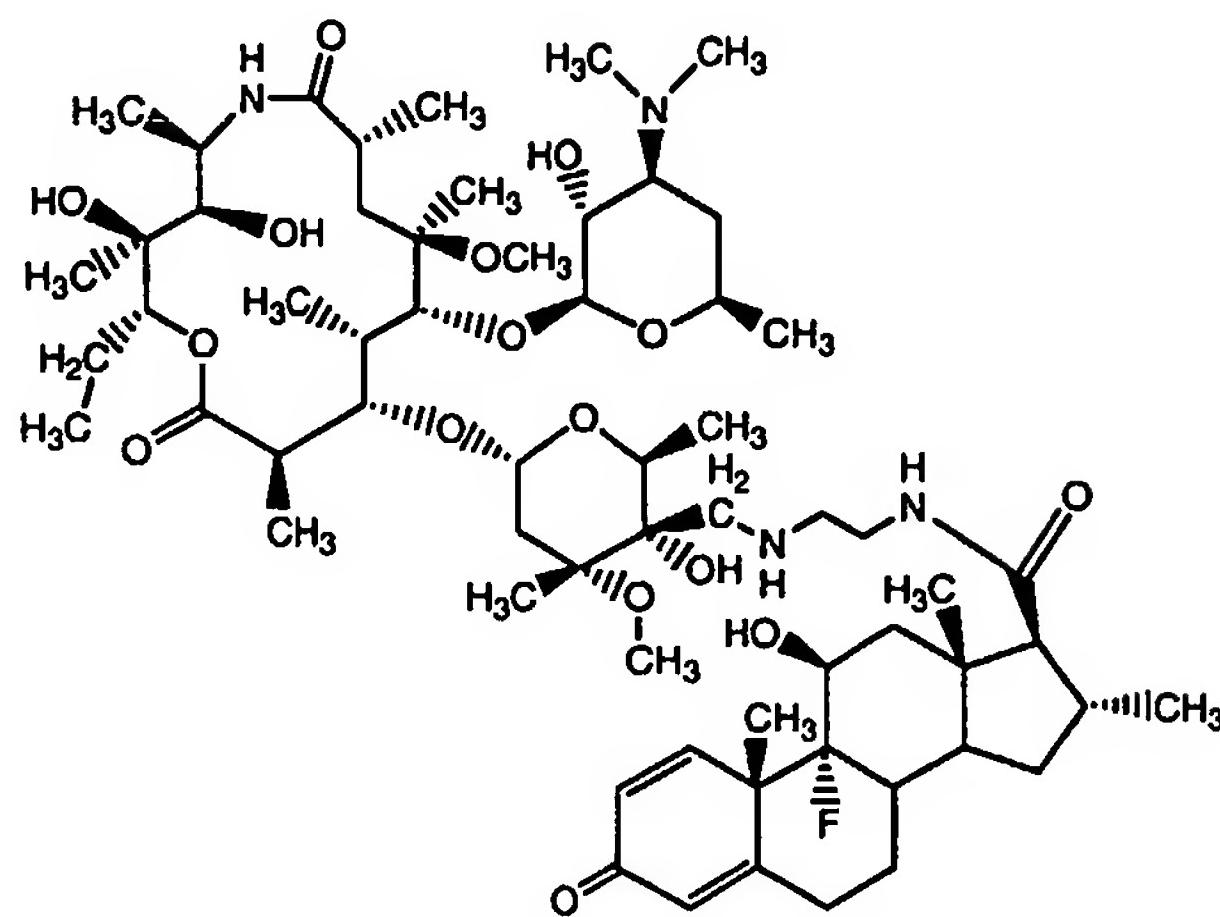


43. A compound 36 characterized by Formula:

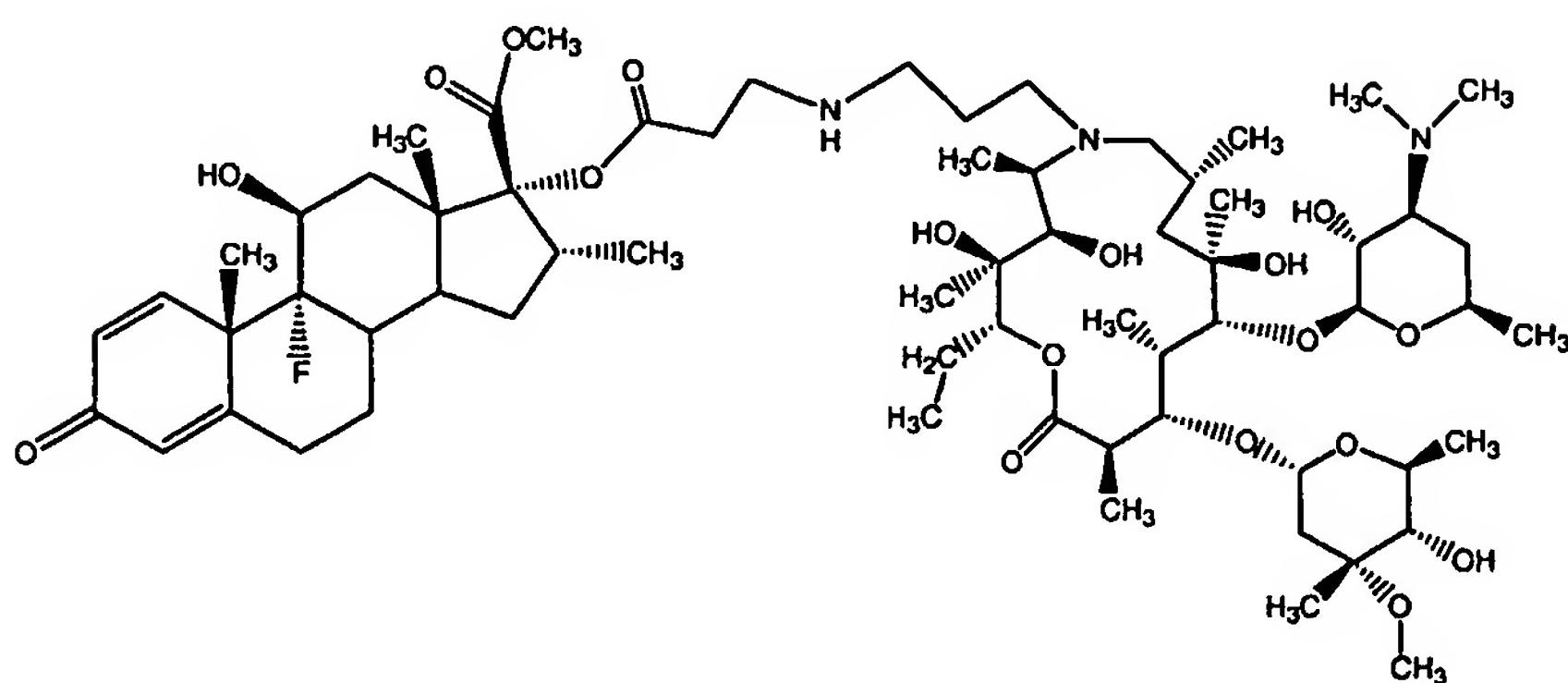


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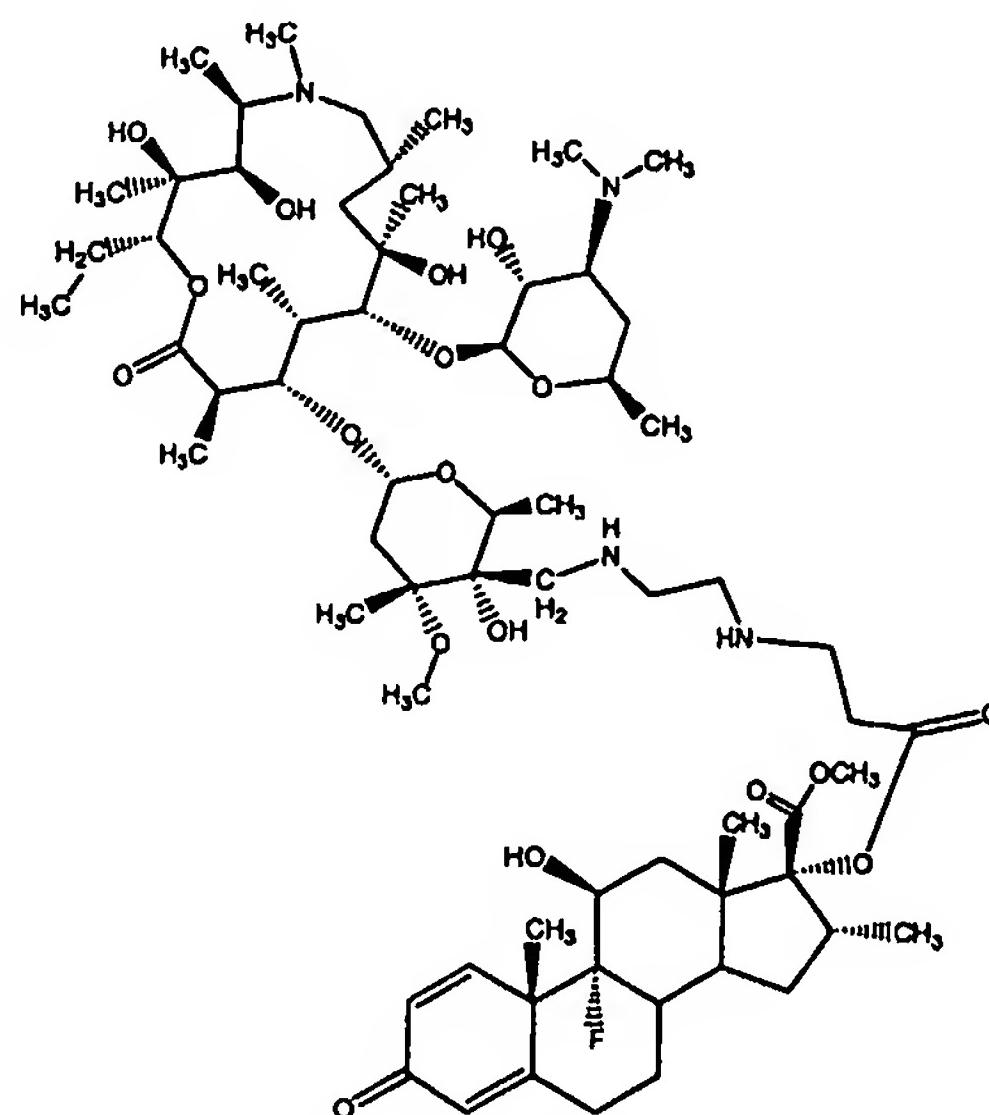
44. A compound 37 characterized by Formula:



45. A compound 38 characterized by Formula:

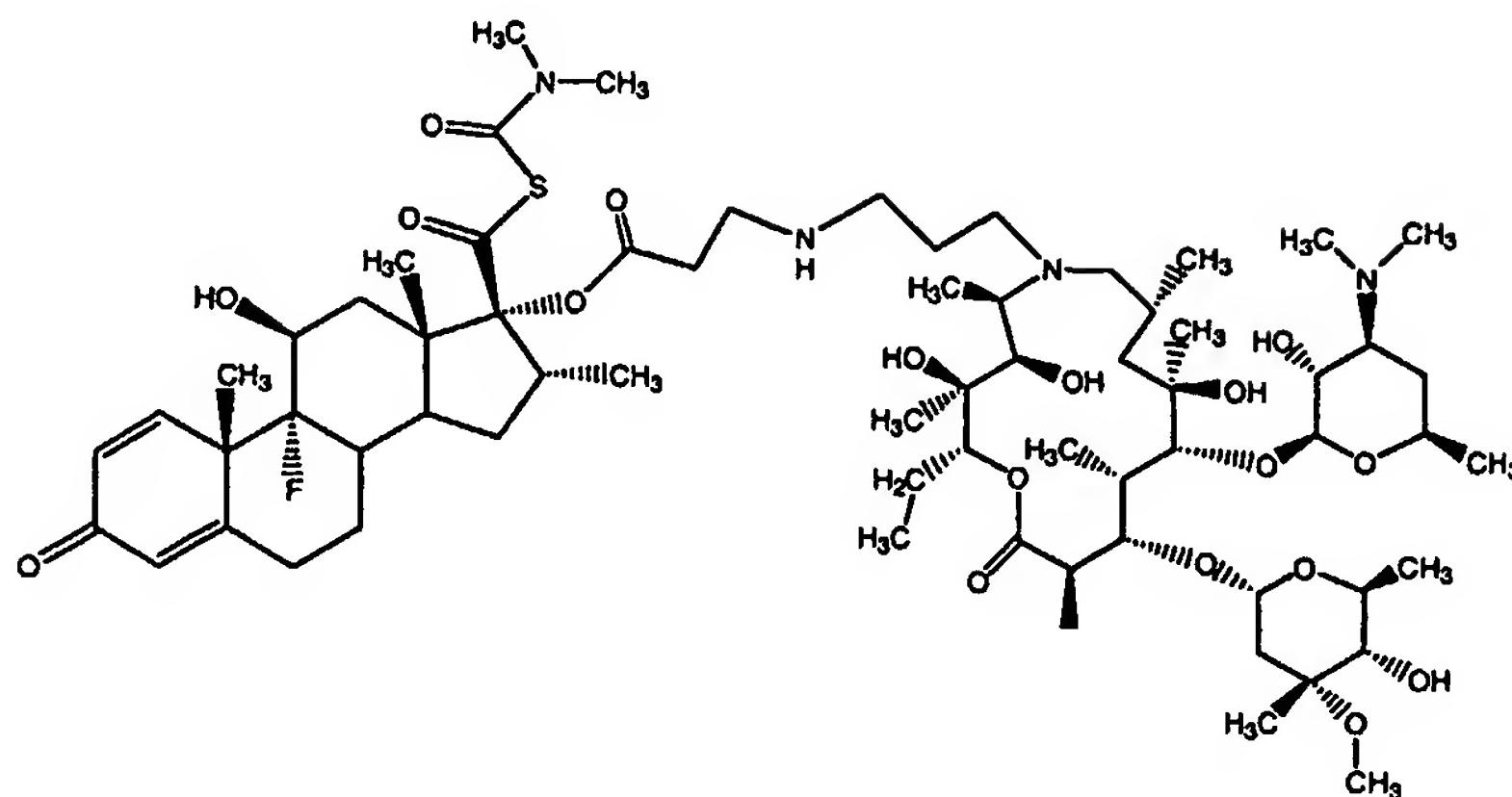


46. A compound 39 characterized by Formula:

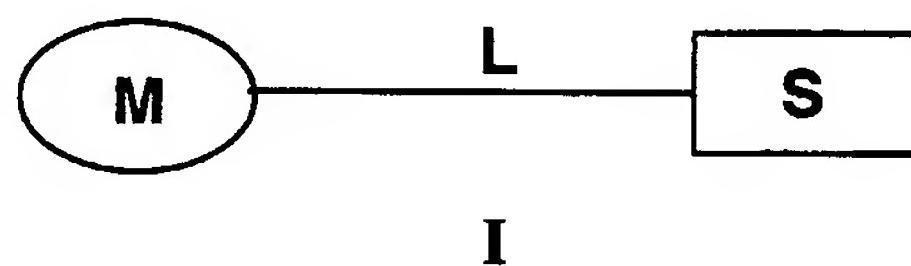


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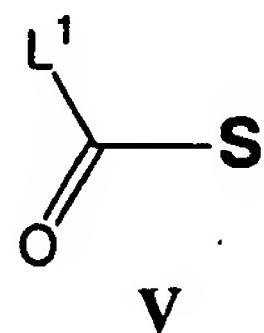
47. A compound 40 characterized by Formula:



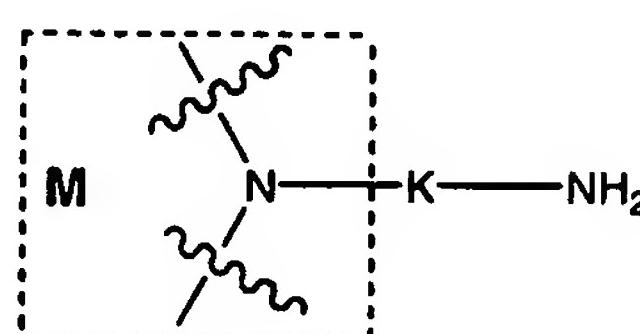
48. A process for the preparation for a compound of Formula I which comprises:



a) for a compound represented by Formula I, where X^2 is $-NHC(O)-$,
by reacting a compound of Formula V:



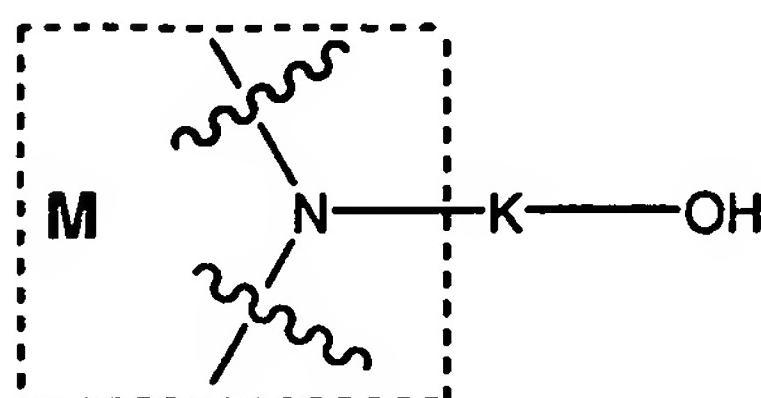
wherein L^1 represents a leaving group, and a free amino group of a macrolide represented by Formula VIId:



VIId

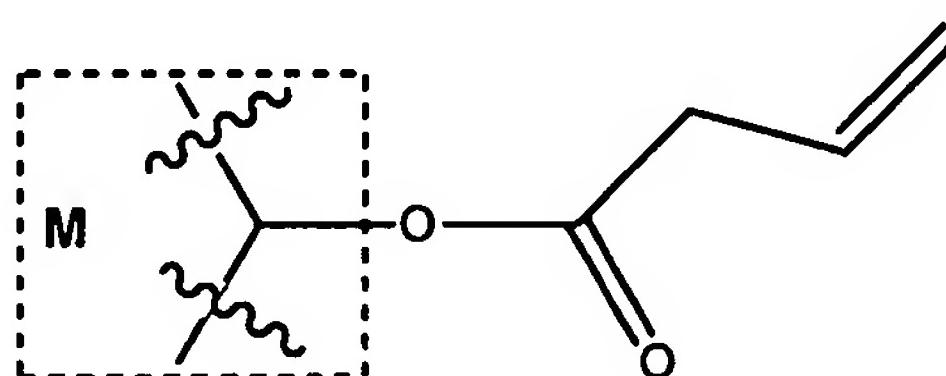
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b) for a compound represented by Formula I, where X^2 is - $OC(O)-$, by reacting a compound of Formula V and a hydroxyl group of a macrolide represented by Formula VIe:

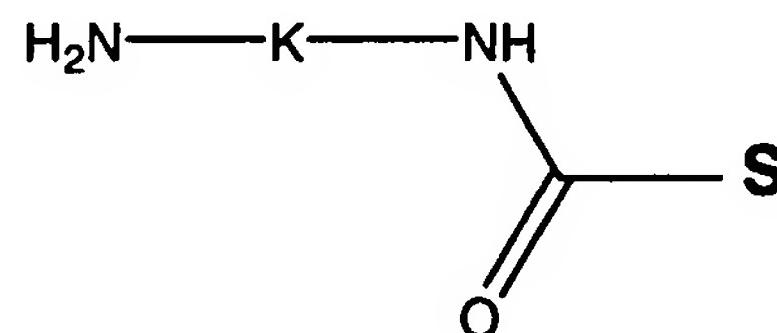


VIe

c) for a compound represented by Formula I, wherein X^1 is - $OC(O)-$, Q is NH and X^2 is - $NHC(O)-$, by reacting a macrolide represented by:

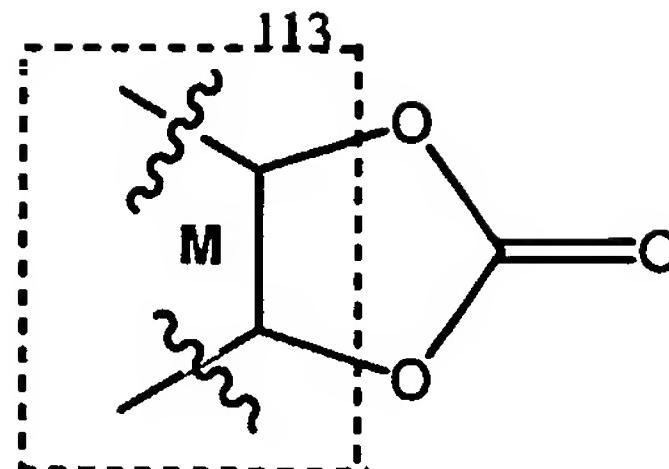


and a free amino group of the compound represented by Formula IVc:

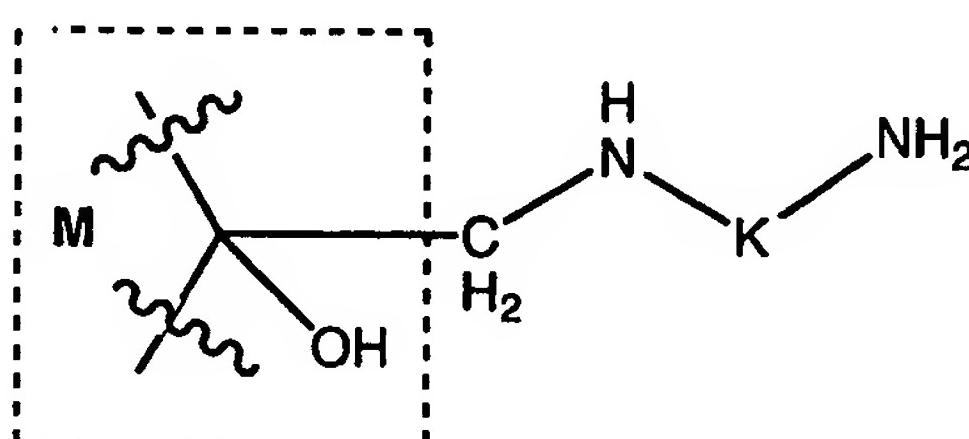


IVc

d) for a compound represented by Formula I, where X^1 is - $OC(O)NH-$ and X^2 is - $NHC(O)-$, by reacting a macrolide represented by Formula VII and a free amino group of Formula IVc:

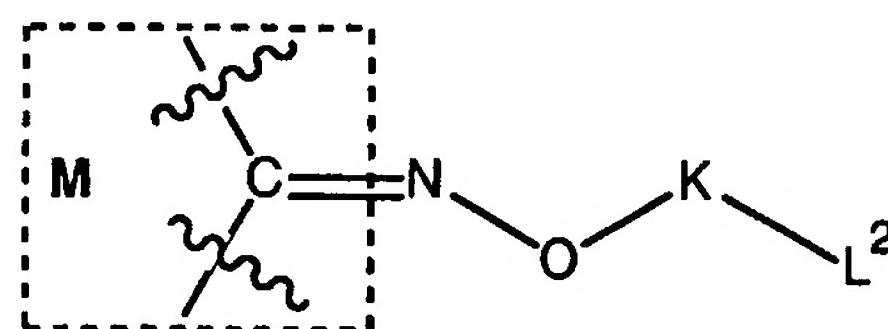


e) for a compound represented by Formula I, where X^1 is $-CH_2-$, Q is $-NH-$ and X^2 is $-NHC(O)-$, by reacting a macrolide represented by Formula Va and a compound of Formula V:

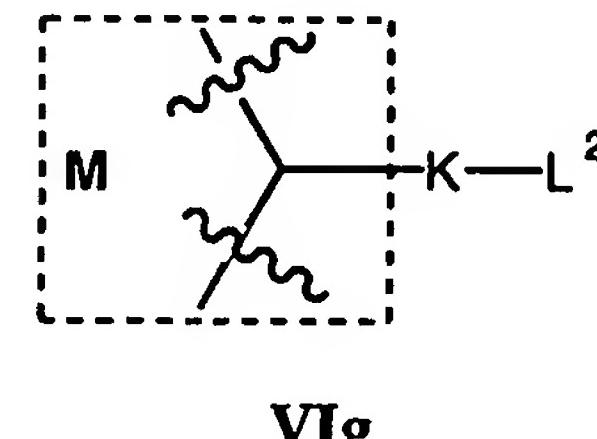
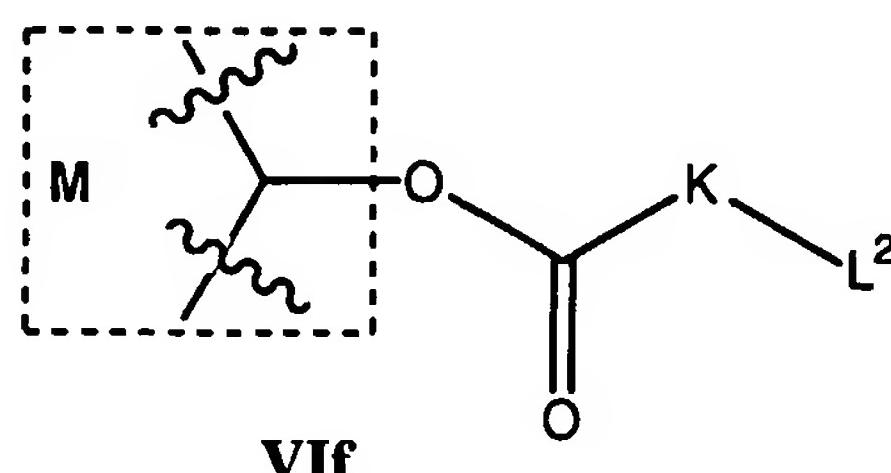


Va

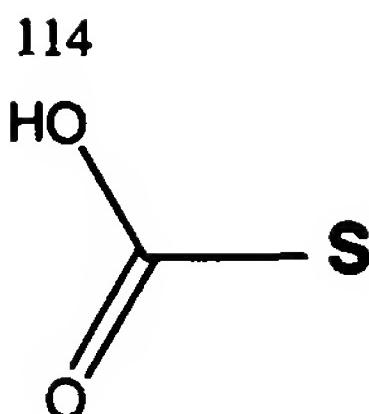
f) compound of Formula I by reacting a macrolide represented by Formula VIIf or by Formula VIg or by Formula VIh having a leaving group L^2



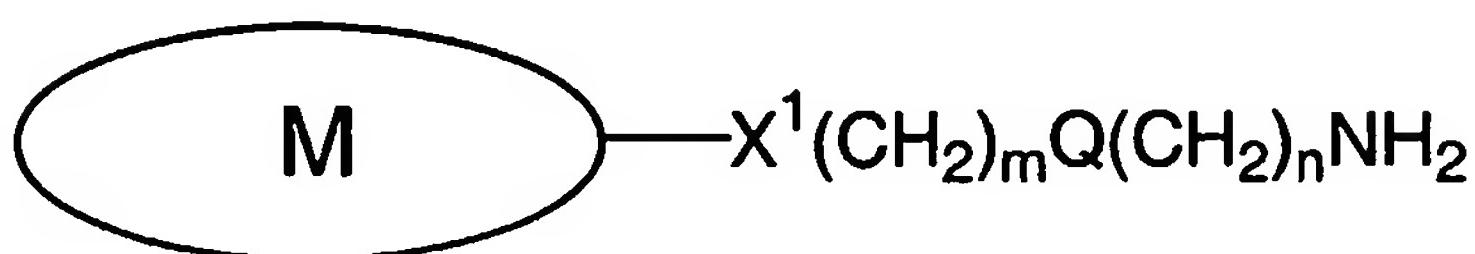
VIh



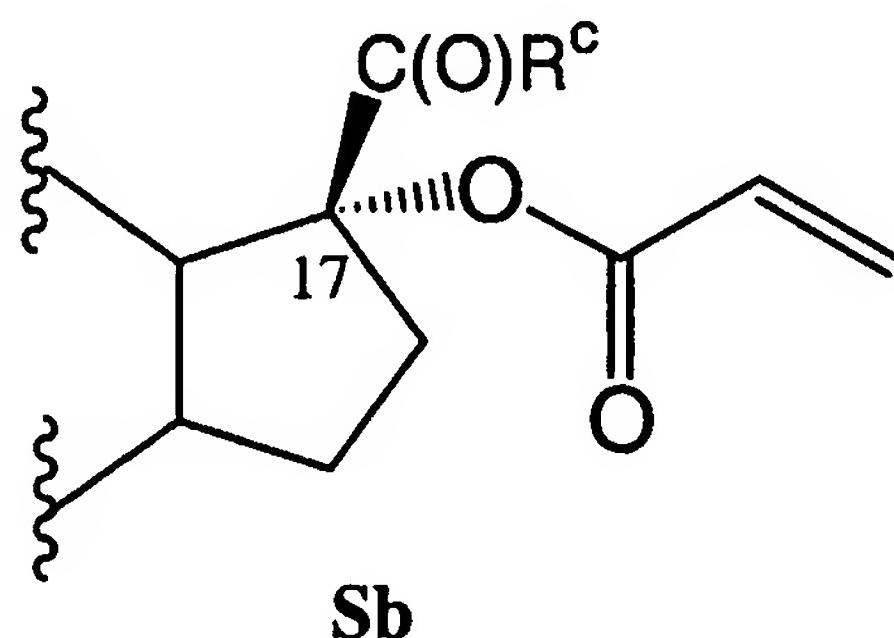
with a free carboxylic acid of steroid represented by Formula IVb.

**IVb**

g) for a compound represented by Formula I, wherein X^1 is $-OC(O)-$, Q is NH and X^2 is $-NH-$ by reacting a macrolide represented by:

**VIId**

and a steroid subunit having a $-C=C-$ bond represented by Formula Sb:



following by modification of R^c group;

49. A pharmaceutical composition comprising a compound as claimed in claims 1 to 46 and pharmaceutically acceptable salts or solvate thereof as well as pharmaceutically acceptable diluent or carrier.

50. The use of compound as claimed in claims 1 to 46 for the manufacture of a medicament for treatment of inflammatory diseases, disorders and conditions characterized by or associated with an undesirable inflammatory immune response,

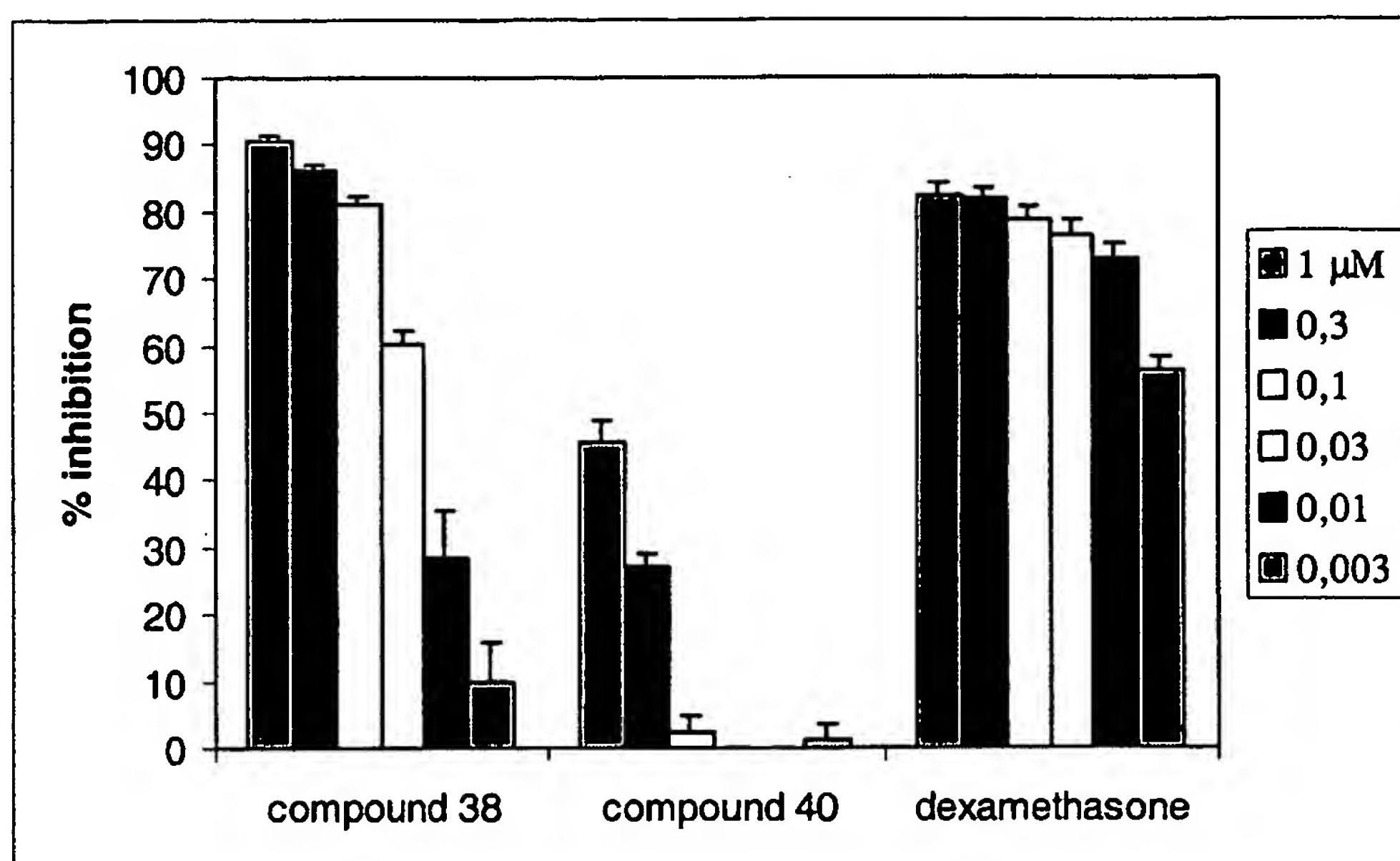
especially of all diseases and conditions induced by or associated with an excessive secretion of TNF- α and IL-1.

51. The use of compound represented by Formula I or a pharmaceutically acceptable salts or solvate thereof for the manufacture of a medicament for treatment of inflammatory conditions and immune or anaphylactic disorders associated with infiltration of leukocytes into inflamed tissue in a subject in need thereof which comprises administering to said subject a therapeutically effective amount of the compound represented by Formula I or a pharmaceutically acceptable salts or solvate thereof.

52. The use of according to claim 50, wherein inflammatory conditions and immune disorders are selected from the group consisting of asthma, adult respiratory distress syndrome, bronchitis, and cystic fibrosis.

53. The use of according to claim 50, wherein said inflammatory conditions and immune disorders are selected from the group consisting of inflammatory conditions or immune disorders of the lungs, joints, eyes, bowel, skin, and heart.

54. The use of according to claim 50, wherein said inflammatory conditions and immune disorders are selected from the group consisting of asthma, adult respiratory distress syndrome, bronchitis, cystic fibrosis, rheumatoid arthritis, rheumatoid spondylitis, osteoarthritis, gouty arthritis, uveitis, conjunctivitis, inflammatory bowel conditions, Crohn's disease, ulcerative colitis, distal proctitis, psoriasis, eczema, dermatitis, coronary infarct damage, chronic inflammation, endotoxin shock, and smooth muscle proliferation disorders.

Figure 1: Inhibition of Hybridoma 13 Proliferation

Inhibition of concanavalin A stimulated interleukin-4 and interleukin-5 production in splenocytes

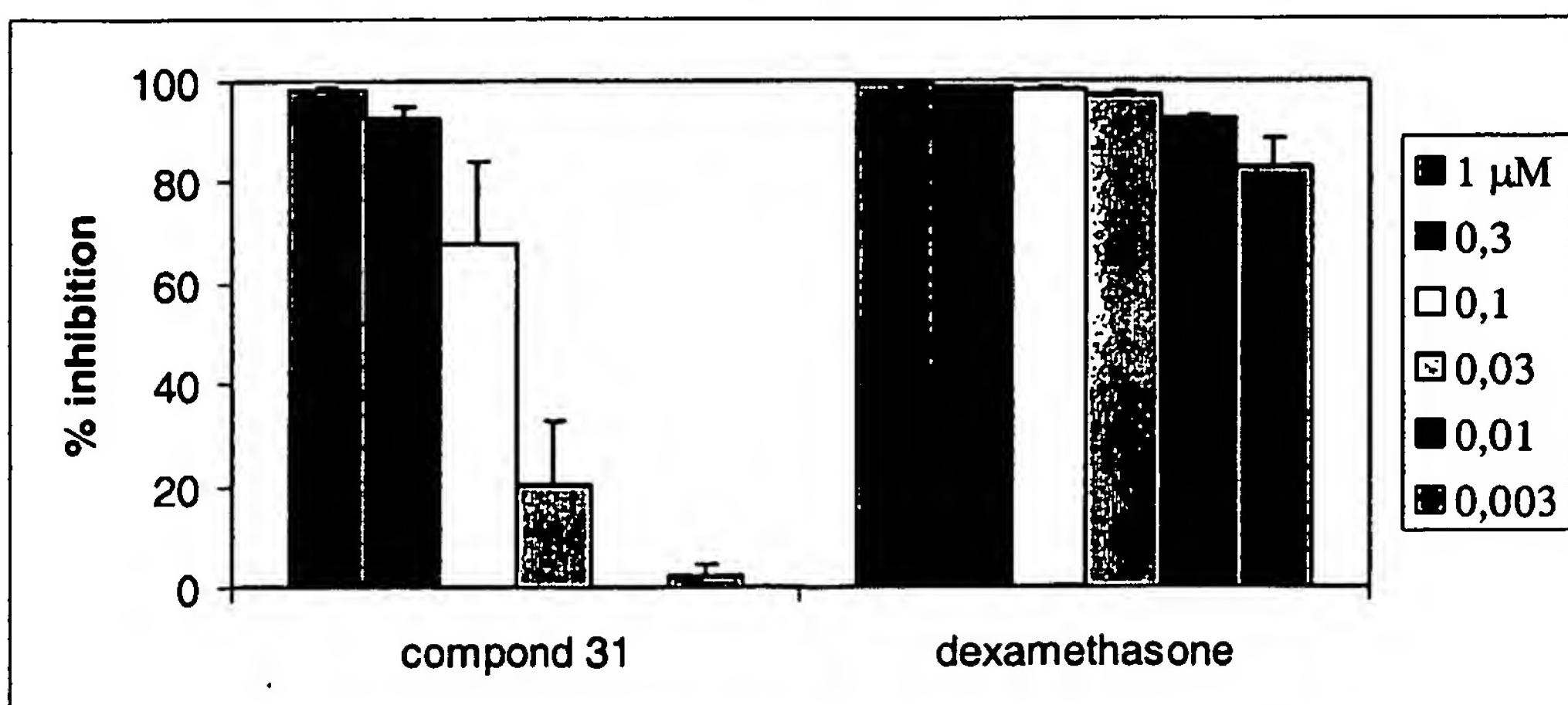
Figure 2: Inhibition of ConA stimulated production of interleukin 4 in murine splenocytes.

Figure 3: Inhibition of ConA stimulated production of interleukin 5 in murine splenocytes

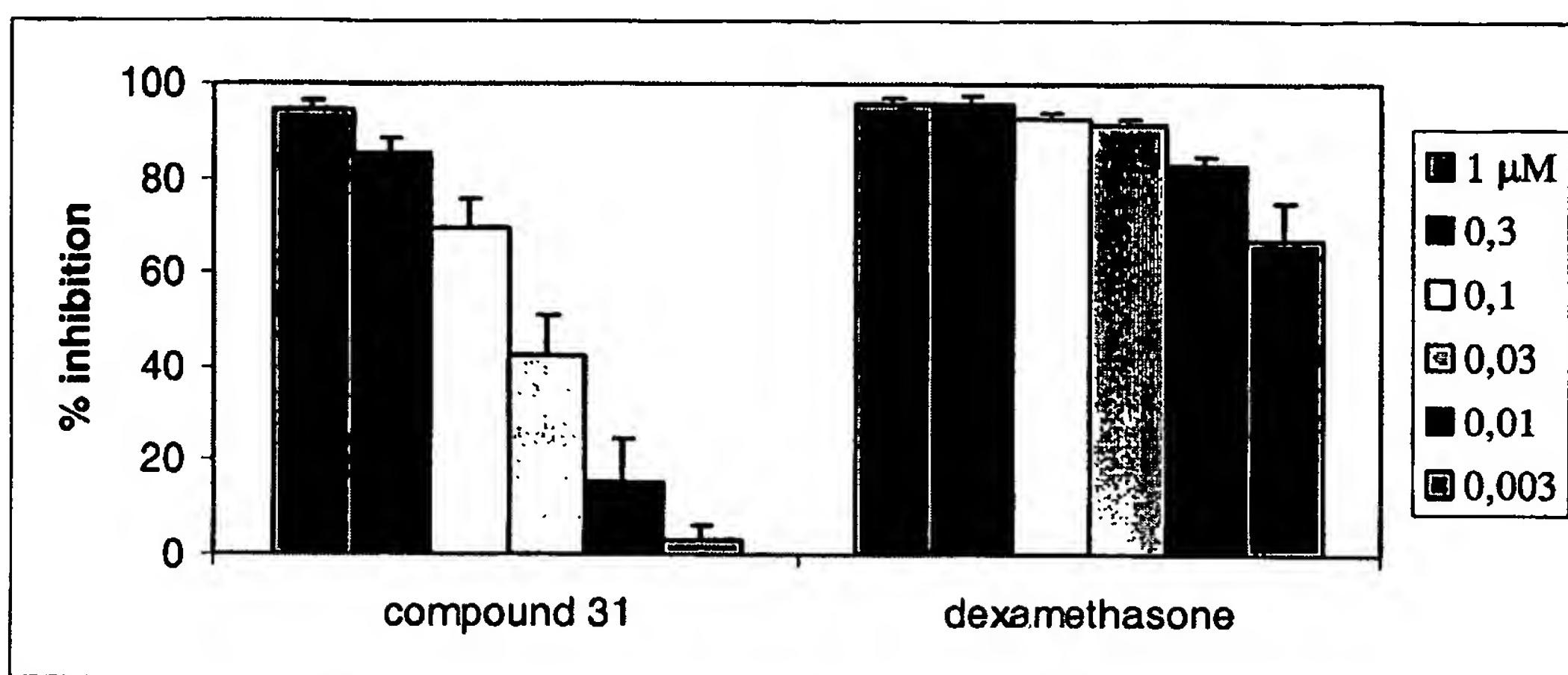
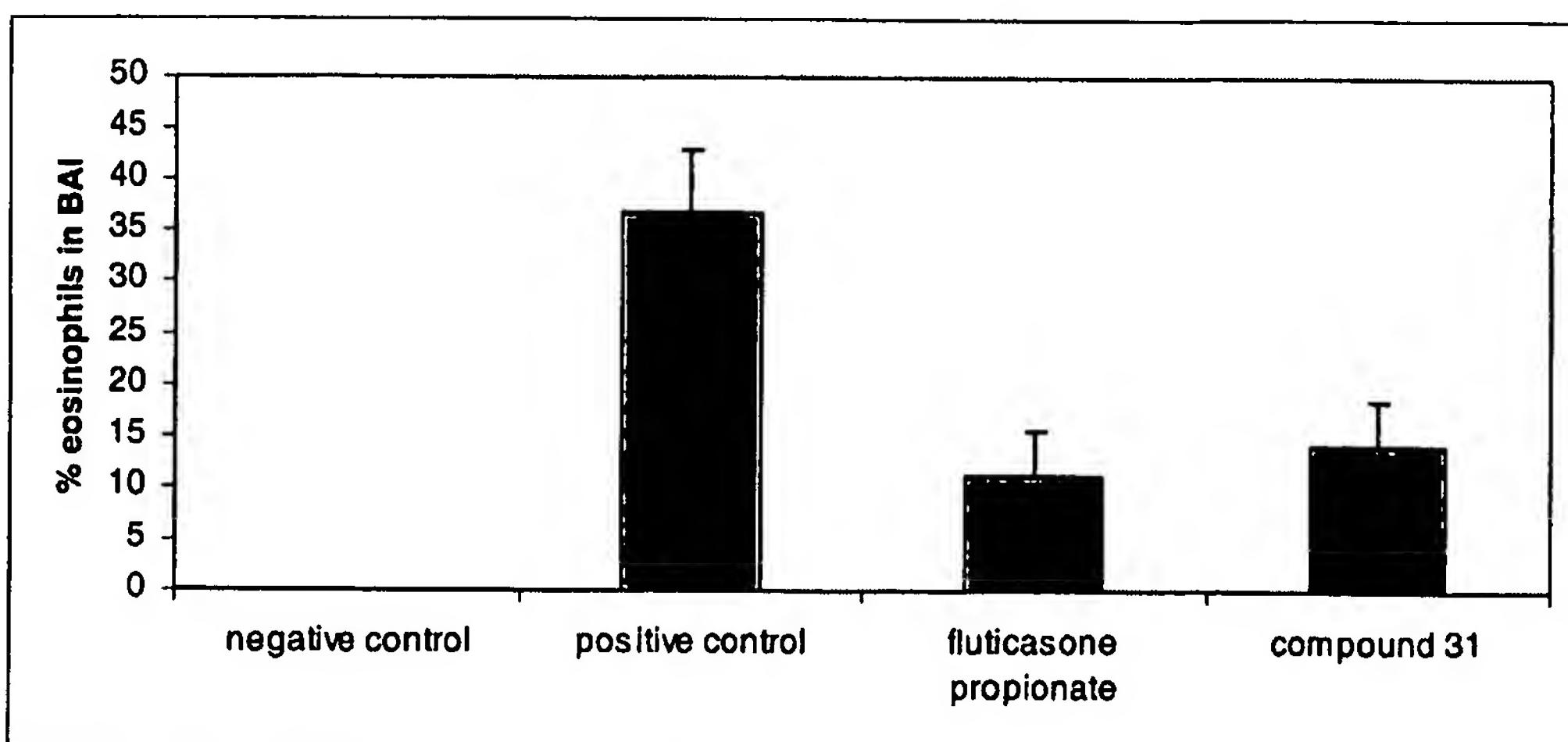


Figure 4: Reduction of eosinophil number in murine model of lung eosinophilia



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